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(54) Title: HAPLOTYPES OF THE EDG6 GENE

(57) Abstract: Novel genetic variants of the Endothelial Differentiation, G-Protein-Coupled Receptor 6 (EDG6) gene are described. Various genotypes, haplotypes, and haplotype pairs that exist in the general United States population are disclosed for the EDG6 gene. Compositions and methods for haplotyping and/or genotyping the EDG6 gene in an individual are also disclosed. Polynucleotides defined by the haplotypes disclosed herein are also described.

HAPLOTYPES OF THE EDG6 GENE

RELATED APPLICATIONS

This application claims the benefit of U.S. Provisional Application Serial No. 60/218,727

5 Serial No. filed July 17, 2000.

FIELD OF THE INVENTION

This invention relates to variation in genes that encode pharmaceutically-important proteins. In particular, this invention provides genetic variants of the human endothelial differentiation, G-10 protein-coupled receptor 6 (EDG6) gene and methods for identifying which variant(s) of this gene is/are possessed by an individual.

BACKGROUND OF THE INVENTION

Current methods for identifying pharmaceuticals to treat disease often start by identifying, 15 cloning, and expressing an important target protein related to the disease. A determination of whether an agonist or antagonist is needed to produce an effect that may benefit a patient with the disease is then made. Then, vast numbers of compounds are screened against the target protein to find new potential drugs. The desired outcome of this process is a lead compound that is specific for the target, thereby reducing the incidence of the undesired side effects usually caused by activity at non-intended 20 targets. The lead compound identified in this screening process then undergoes further *in vitro* and *in vivo* testing to determine its absorption, disposition, metabolism and toxicological profiles. Typically, this testing involves use of cell lines and animal models with limited, if any, genetic diversity.

What this approach fails to consider, however, is that natural genetic variability exists between individuals in any and every population with respect to pharmaceutically-important proteins, including 25 the protein targets of candidate drugs, the enzymes that metabolize these drugs and the proteins whose activity is modulated by such drug targets. Subtle alteration(s) in the primary nucleotide sequence of a gene encoding a pharmaceutically-important protein may be manifested as significant variation in expression, structure and/or function of the protein. Such alterations may explain the relatively high degree of uncertainty inherent in the treatment of individuals with a drug whose design is based upon a 30 single representative example of the target or enzyme(s) involved in metabolizing the drug. For example, it is well-established that some drugs frequently have lower efficacy in some individuals than others, which means such individuals and their physicians must weigh the possible benefit of a larger dosage against a greater risk of side effects. Also, there is significant variation in how well people metabolize drugs and other exogenous chemicals, resulting in substantial interindividual variation in 35 the toxicity and/or efficacy of such exogenous substances (Evans et al., 1999, *Science* 286:487-491). This variability in efficacy or toxicity of a drug in genetically-diverse patients makes many drugs ineffective or even dangerous in certain groups of the population, leading to the failure of such drugs in clinical trials or their early withdrawal from the market even though they could be highly beneficial for

other groups in the population. This problem significantly increases the time and cost of drug discovery and development, which is a matter of great public concern.

It is well-recognized by pharmaceutical scientists that considering the impact of the genetic variability of pharmaceutically-important proteins in the early phases of drug discovery and development is likely to reduce the failure rate of candidate and approved drugs (Marshall A 1997 *Nature Biotech* **15**:1249-52; Kleyn PW et al. 1998 *Science* **281**: 1820-21; Kola I 1999 *Curr Opin Biotech* **10**:589-92; Hill AVS et al. 1999 in *Evolution in Health and Disease* Stearns SS (Ed.) Oxford University Press, New York, pp 62-76; Meyer U.A. 1999 in *Evolution in Health and Disease* Stearns SS (Ed.) Oxford University Press, New York, pp 41-49; Kalow W et al. 1999 *Clin. Pharm. Therap.* **66**:445-7; Marshall, E 1999 *Science* **284**:406-7; Judson R et al. 2000 *Pharmacogenomics* **1**:1-12; Roses AD 2000 *Nature* **405**:857-65). However, in practice this has been difficult to do, in large part because of the time and cost required for discovering the amount of genetic variation that exists in the population (Chakravarti A 1998 *Nature Genet* **19**:216-7; Wang DG et al 1998 *Science* **280**:1077-82; Chakravarti A 1999 *Nat Genet* **21**:56-60 (suppl); Stephens JC 1999 *Mol. Diagnosis* **4**:309-317; Kwok PY and Gu S 1999 *Mol. Med. Today* **5**:538-43; Davidson S 2000 *Nature Biotech* **18**:1134-5).

The standard for measuring genetic variation among individuals is the haplotype, which is the ordered combination of polymorphisms in the sequence of each form of a gene that exists in the population. Because haplotypes represent the variation across each form of a gene, they provide a more accurate and reliable measurement of genetic variation than individual polymorphisms. For example, while specific variations in gene sequences have been associated with a particular phenotype such as disease susceptibility (Roses AD *supra*; Ulbrecht M et al. 2000 *Am J Respir Crit Care Med* **161**: 469-74) and drug response (Wolfe CR et al. 2000 *BMJ* **320**:987-90; Dahl BS 1997 *Acta Psychiatr Scand* **96** (Suppl 391): 14-21), in many other cases an individual polymorphism may be found in a variety of genomic backgrounds, i.e., different haplotypes, and therefore shows no definitive coupling between the polymorphism and the causative site for the phenotype (Clark AG et al. 1998 *Am J Hum Genet* **63**:595-612; Ulbrecht M et al. 2000 *supra*; Drysdale et al. 2000 *PNAS* **97**:10483-10488). Thus, there is an unmet need in the pharmaceutical industry for information on what haplotypes exist in the population for pharmaceutically-important genes. Such haplotype information would be useful in improving the efficiency and output of several steps in the drug discovery and development process, including target validation, identifying lead compounds, and early phase clinical trials (Marshall et al., *supra*).

One pharmaceutically-important gene for the treatment of cancer, angiogenesis and inflammation is the endothelial differentiation, G-protein-coupled receptor 6 (EDG6) gene or its encoded product. EDG receptors, such as EDG6, constitute a novel subfamily of G-protein-coupled receptors displaying a heterogeneous expression pattern. Members of this family can bind lysophospholipids or lysosphingolipids as ligands. EDG6 is specifically expressed in fetal and adult lymphoid and hematopoietic tissue as well as in lung (Graler et al., *Genomics* 1998; **53**:164-169). Graler et al. (*supra*) suggest that because of the known mitogenic and chemotactic activity of bioactive

lipids, EDG6 may play an essential role in lymphocyte cell signaling. EDG6 can also bind sphingosine 1-phosphate, a lysolipid, to elicit biological responses, including mitogenesis, differentiation, migration and apoptosis, via receptor-dependent mechanisms. Sphingosine 1-phosphate has been implicated in pathophysiological disease states, such as cancer, angiogenesis and

5 inflammation (Pyne and Pyne, *Biochem J* 2000; 349:385-402). For example, sphingosine 1-phosphate (S1-P) has been shown to induce the secretion of type H Insulin-like growth factor II, which is responsible for proliferation of cultured breast cancer cells (Goetzel et al., *Cancer Res.* 1999; 59:4732-4737). Goetzel et al. have shown that another EDG receptor, EDG4, is a marker for ovarian cancer, and it is possible that other S1-P-specific EDG receptors may be involved in cancer. Therefore, aberrant

10 expression of EDG6 may result in changes in S1-P concentrations, which could affect several disease processes.

The endothelial differentiation, G-protein-coupled receptor 6 gene is located on chromosome 19p13.3 and contains 1 exon that encodes a 384 amino acid protein. A reference sequence for the EDG6 gene is shown in the contiguous lines of Figure 1 (Genaissance Reference No. 3216828; SEQ 15 ID NO: 1). Reference sequences for the coding sequence (GenBank Accession No. NM_003775.1) and protein are shown in Figures 2 (SEQ ID NO: 2) and 3 (SEQ ID NO: 3), respectively.

Because of the potential for variation in the EDG6 gene to affect the expression and function of the encoded protein, it would be useful to know whether polymorphisms exist in the EDG6 gene, as well as how such polymorphisms are combined in different copies of the gene. Such information 20 could be applied for studying the biological function of EDG6 as well as in identifying drugs targeting this protein for the treatment of disorders related to its abnormal expression or function.

SUMMARY OF THE INVENTION

Accordingly, the inventors herein have discovered 23 novel polymorphic sites in the EDG6 25 gene. These polymorphic sites (PS) correspond to the following nucleotide positions in Figure 1: 3591 (PS1), 3697 (PS2), 3804 (PS3), 3818 (PS4), 4123 (PS5), 4240 (PS6), 4472 (PS7), 4499 (PS8), 4531 (PS9), 4574 (PS10), 4736 (PS11), 4813 (PS12), 5068 (PS13), 5103 (PS14), 5150 (PS15), 5179 (PS16), 5301 (PS17), 5333 (PS18), 5448 (PS19), 5560 (PS20), 5580 (PS21), 5587 (PS22) and 5606 (PS23). The polymorphisms at these sites are guanine or adenine at PS1, cytosine or thymine at PS2, 30 cytosine or thymine at PS3, adenine or guanine at PS4, cytosine or thymine at PS5, guanine or adenine at PS6, guanine or adenine at PS7, guanine or adenine at PS8, guanine or adenine at PS9, guanine or thymine at PS10, cytosine or thymine at PS11, cytosine or thymine at PS12, cytosine or thymine at PS13, guanine or thymine at PS14, guanine or adenine at PS15, guanine or adenine at PS16, guanine or adenine at PS17, guanine or adenine at PS18, guanine or cytosine at PS19, guanine or adenine at PS20, 35 guanine or adenine at PS21, cytosine or thymine at PS22 and guanine or cytosine at PS23. In addition, the inventors have determined the identity of the alleles at these sites in a human reference population of 79 unrelated individuals self-identified as belonging to one of four major population groups: African descent, Asian, Caucasian and Hispanic/Latino. From this information, the inventors deduced a set of

haplotypes and haplotype pairs for PS1-PS23 in the EDG6 gene, which are shown below in Tables 5 and 4, respectively. Each of these EDG6 haplotypes defines a naturally-occurring isoform (also referred to herein as an “isogene”) of the EDG6 gene that exists in the human population. The frequency with which each haplotype and haplotype pair occurs within the total reference population 5 and within each of the four major population groups included in the reference population was also determined.

Thus, in one embodiment, the invention provides a method, composition and kit for genotyping the EDG6 gene in an individual. The genotyping method comprises identifying the nucleotide pair that is present at one or more polymorphic sites selected from the group consisting of 10 PS1, PS2, PS3, PS4, PS5, PS6, PS7, PS8, PS9, PS10, PS11, PS12, PS13, PS14, PS15, PS16, PS17, PS18, PS19, PS20, PS21, PS22 and PS23 in both copies of the EDG6 gene from the individual. A genotyping composition of the invention comprises an oligonucleotide probe or primer which is designed to specifically hybridize to a target region containing, or adjacent to, one of these novel 15 EDG6 polymorphic sites. A genotyping kit of the invention comprises a set of oligonucleotides designed to genotype each of these novel EDG6 polymorphic sites. The genotyping method, composition, and kit are useful in determining whether an individual has one of the haplotypes in 20 Table 5 below or has one of the haplotype pairs in Table 4 below.

The invention also provides a method for haplotyping the EDG6 gene in an individual. In one embodiment, the haplotyping method comprises determining, for one copy of the EDG6 gene, the 25 identity of the nucleotide at one or more polymorphic sites selected from the group consisting of PS1, PS2, PS3, PS4, PS5, PS6, PS7, PS8, PS9, PS10, PS11, PS12, PS13, PS14, PS15, PS16, PS17, PS18, PS19, PS20, PS21, PS22 and PS23. In another embodiment, the haplotyping method comprises determining whether one copy of the individual’s EDG6 gene is defined by one of the EDG6 30 haplotypes shown in Table 5, below, or a sub-haplotype thereof. In a preferred embodiment, the haplotyping method comprises determining whether both copies of the individual’s EDG6 gene are defined by one of the EDG6 haplotype pairs shown in Table 4 below, or a sub-haplotype pair thereof. The method for establishing the EDG6 haplotype or haplotype pair of an individual is useful for improving the efficiency and reliability of several steps in the discovery and development of drugs for 35 treating diseases associated with EDG6 activity, e.g., cancer, angiogenesis and inflammation.

For example, the haplotyping method can be used by the pharmaceutical research scientist to validate EDG6 as a candidate target for treating a specific condition or disease predicted to be associated with EDG6 activity. Determining for a particular population the frequency of one or more of the individual EDG6 haplotypes or haplotype pairs described herein will facilitate a decision on whether to pursue EDG6 as a target for treating the specific disease of interest. In particular, if 35 variable EDG6 activity is associated with the disease, then one or more EDG6 haplotypes or haplotype pairs will be found at a higher frequency in disease cohorts than in appropriately genetically matched controls. Conversely, if each of the observed EDG6 haplotypes are of similar frequencies in the disease and control groups, then it may be inferred that variable EDG6 activity has little, if any,

involvement with that disease. In either case, the pharmaceutical research scientist can, without *a priori* knowledge as to the phenotypic effect of any EDG6 haplotype or haplotype pair, apply the information derived from detecting EDG6 haplotypes in an individual to decide whether modulating EDG6 activity would be useful in treating the disease.

5 The claimed invention is also useful in screening for compounds targeting EDG6 to treat a specific condition or disease predicted to be associated with EDG6 activity. For example, detecting which of the EDG6 haplotypes or haplotype pairs disclosed herein are present in individual members of a population with the specific disease of interest enables the pharmaceutical scientist to screen for a compound(s) that displays the highest desired agonist or antagonist activity for each of the most frequent EDG6 isoforms present in the disease population. Thus, without requiring any *a priori* knowledge of the phenotypic effect of any particular EDG6 haplotype or haplotype pair, the claimed haplotyping method provides the scientist with a tool to identify lead compounds that are more likely to show efficacy in clinical trials.

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15 The method for haplotyping the EDG6 gene in an individual is also useful in the design of clinical trials of candidate drugs for treating a specific condition or disease predicted to be associated with EDG6 activity. For example, instead of randomly assigning patients with the disease of interest to the treatment or control group as is typically done now, determining which of the EDG6 haplotype(s) disclosed herein are present in individual patients enables the pharmaceutical scientist to distribute EDG6 haplotypes and/or haplotype pairs evenly to treatment and control groups, thereby

20 reducing the potential for bias in the results that could be introduced by a larger frequency of an EDG6 haplotype or haplotype pair that had a previously unknown association with response to the drug being studied in the trial. Thus, by practicing the claimed invention, the scientist can more confidently rely on the information learned from the trial, without first determining the phenotypic effect of any EDG6 haplotype or haplotype pair.

25 In another embodiment, the invention provides a method for identifying an association between a trait and an EDG6 genotype, haplotype, or haplotype pair for one or more of the novel polymorphic sites described herein. The method comprises comparing the frequency of the EDG6 genotype, haplotype, or haplotype pair in a population exhibiting the trait with the frequency of the EDG6 genotype or haplotype in a reference population. A higher frequency of the EDG6 genotype, haplotype, or haplotype pair in the trait population than in the reference population indicates the trait is associated with the EDG6 genotype, haplotype, or haplotype pair. In preferred embodiments, the trait is susceptibility to a disease, severity of a disease, the staging of a disease or response to a drug. In a particularly preferred embodiment, the EDG6 haplotype is selected from the haplotypes shown in Table 5, or a sub-haplotype thereof. Such methods have applicability in developing diagnostic tests and therapeutic treatments for cancer, angiogenesis and inflammation.

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In yet another embodiment, the invention provides an isolated polynucleotide comprising a nucleotide sequence which is a polymorphic variant of a reference sequence for the EDG6 gene or a fragment thereof. The reference sequence comprises the contiguous sequences shown in Figure 1 and

the polymorphic variant comprises at least one polymorphism selected from the group consisting of adenine at PS1, thymine at PS2, thymine at PS3, guanine at PS4, thymine at PS5, adenine at PS6, adenine at PS7, adenine at PS8, adenine at PS9, thymine at PS10, thymine at PS11, thymine at PS12, thymine at PS13, thymine at PS14, adenine at PS15, adenine at PS16, adenine at PS17, adenine at 5 PS18, cytosine at PS19, adenine at PS20, adenine at PS21, thymine at PS22 and cytosine at PS23.

A particularly preferred polymorphic variant is an isogene of the EDG6 gene. An EDG6 isogene of the invention comprises guanine or adenine at PS1, cytosine or thymine at PS2, cytosine or thymine at PS3, adenine or guanine at PS4, cytosine or thymine at PS5, guanine or adenine at PS6, guanine or adenine at PS7, guanine or adenine at PS8, guanine or adenine at PS9, guanine or thymine 10 at PS10, cytosine or thymine at PS11, cytosine or thymine at PS12, cytosine or thymine at PS13, guanine or thymine at PS14, guanine or adenine at PS15, guanine or adenine at PS16, guanine or adenine at PS17, guanine or adenine at PS18, guanine or cytosine at PS19, guanine or adenine at PS20, guanine or adenine at PS21, cytosine or thymine at PS22 and guanine or cytosine at PS23. The invention also provides a collection of EDG6 isogenes, referred to herein as an EDG6 genome 15 anthology.

In another embodiment, the invention provides a polynucleotide comprising a polymorphic variant of a reference sequence for an EDG6 cDNA or a fragment thereof. The reference sequence comprises SEQ ID NO:2 (Fig.2) and the polymorphic cDNA comprises at least one polymorphism selected from the group consisting of thymine at a position corresponding to nucleotide 114, adenine at 20 a position corresponding to nucleotide 231, adenine at a position corresponding to nucleotide 463, adenine at a position corresponding to nucleotide 490, adenine at a position corresponding to nucleotide 522, thymine at a position corresponding to nucleotide 565, thymine at a position corresponding to nucleotide 727, thymine at a position corresponding to nucleotide 804, thymine at a position corresponding to nucleotide 1059, thymine at a position corresponding to nucleotide 1094 and 25 adenine at a position corresponding to nucleotide 1141. A particularly preferred polymorphic cDNA variant comprises the coding sequence of an EDG6 isogene defined by haplotypes 3c, 7c-12c, 19c-22c, and 24c.

Polynucleotides complementary to these EDG6 genomic and cDNA variants are also provided by the invention. It is believed that polymorphic variants of the EDG6 gene will be useful in studying 30 the expression and function of EDG6, and in expressing EDG6 protein for use in screening for candidate drugs to treat diseases related to EDG6 activity.

In other embodiments, the invention provides a recombinant expression vector comprising one of the polymorphic genomic variants operably linked to expression regulatory elements as well as a recombinant host cell transformed or transfected with the expression vector. The recombinant vector 35 and host cell may be used to express EDG6 for protein structure analysis and drug binding studies.

In yet another embodiment, the invention provides a polypeptide comprising a polymorphic variant of a reference amino acid sequence for the EDG6 protein. The reference amino acid sequence comprises SEQ ID NO:3 (Fig.3) and the polymorphic variant comprises at least one variant amino acid

selected from the group consisting of arginine at a position corresponding to amino acid position 155, serine at a position corresponding to amino acid position 164, serine at a position corresponding to amino acid position 189, cysteine at a position corresponding to amino acid position 243, leucine at a position corresponding to amino acid position 365 and methionine at a position corresponding to amino acid position 381. A polymorphic variant of EDG6 is useful in studying the effect of the variation on the biological activity of EDG6 as well as on the binding affinity of candidate drugs targeting EDG6 for the treatment of cancer, angiogenesis and inflammation.

The present invention also provides antibodies that recognize and bind to the above polymorphic EDG6 protein variant. Such antibodies can be utilized in a variety of diagnostic and prognostic formats and therapeutic methods.

The present invention also provides nonhuman transgenic animals comprising one of the EDG6 polymorphic genomic variants described herein and methods for producing such animals. The transgenic animals are useful for studying expression of the EDG6 isogenes *in vivo*, for *in vivo* screening and testing of drugs targeted against EDG6 protein, and for testing the efficacy of therapeutic agents and compounds for cancer, angiogenesis and inflammation in a biological system.

The present invention also provides a computer system for storing and displaying polymorphism data determined for the EDG6 gene. The computer system comprises a computer processing unit; a display; and a database containing the polymorphism data. The polymorphism data includes the polymorphisms, the genotypes and the haplotypes identified for the EDG6 gene in a reference population. In a preferred embodiment, the computer system is capable of producing a display showing EDG6 haplotypes organized according to their evolutionary relationships.

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 illustrates a reference sequence for the EDG6 gene (Genaissance Reference No. 3216828; contiguous lines), with the start and stop positions of each region of coding sequence indicated with a bracket ([or]) and the numerical position below the sequence and the polymorphic site(s) and polymorphism(s) identified by Applicants in a reference population indicated by the variant nucleotide positioned below the polymorphic site in the sequence. SEQ ID NO:1 is equivalent to Figure 1, with the two alternative allelic variants of each polymorphic site indicated by the appropriate nucleotide symbol (R= G or A, Y= T or C, M= A or C, K= G or T, S= G or C, and W= A or T; WIPO standard ST.25). SEQ ID NO:119 is a modified version of SEQ ID NO:1 that shows the context sequence of each polymorphic site, PS1-PS23, in a uniform format to facilitate electronic searching. For each polymorphic site, SEQ ID NO:119 contains a block of 60 bases of the nucleotide sequence encompassing the centrally-located polymorphic site at the 30th position, followed by 60 bases of unspecified sequence to represent that each PS is separated by genomic sequence whose composition is defined elsewhere herein.

Figure 2 illustrates a reference sequence for the EDG6 coding sequence (contiguous lines; SEQ ID NO:2), with the polymorphic site(s) and polymorphism(s) identified by Applicants in a

reference population indicated by the variant nucleotide positioned below the polymorphic site in the sequence.

Figure 3 illustrates a reference sequence for the EDG6 protein (contiguous lines; SEQ ID NO:3), with the variant amino acid(s) caused by the polymorphism(s) of Figure 2 positioned below the 5 polymorphic site in the sequence.

DESCRIPTION OF THE PREFERRED EMBODIMENTS

The present invention is based on the discovery of novel variants of the EDG6 gene. As described in more detail below, the inventors herein discovered 24 isogenes of the EDG6 gene by 10 characterizing the EDG6 gene found in genomic DNAs isolated from an Index Repository that contains immortalized cell lines from one chimpanzee and 93 human individuals. The human individuals included a reference population of 79 unrelated individuals self-identified as belonging to one of four major population groups: Caucasian (21 individuals), African descent (20 individuals), Asian (20 individuals), or Hispanic/Latino (18 individuals). To the extent possible, the members of 15 this reference population were organized into population subgroups by their self-identified ethnogeographic origin as shown in Table 1 below.

Table 1. Population Groups in the Index Repository

Population Group	Population Subgroup	No. of Individuals
African descent		20
	Sierra Leone	1
Asian		20
	Burma	1
	China	3
	Japan	6
	Korea	1
	Philippines	5
	Vietnam	4
Caucasian		21
	British Isles	3
	British Isles/Central	4
	British Isles/Eastern	1
	Central/Eastern	1
	Eastern	3
	Central/Mediterranean	1
	Mediterranean	2
	Scandinavian	2
Hispanic/Latino		18
	Caribbean	8
	Caribbean (Spanish Descent)	2
	Central American (Spanish Descent)	1
	Mexican American	4
	South American (Spanish Descent)	3

20 In addition, the Index Repository contains three unrelated indigenous American Indians (one from each of North, Central and South America), one three-generation Caucasian family (from the

CEPH Utah cohort) and one two-generation African-American family.

The EDG6 isogenes present in the human reference population are defined by haplotypes for 23 polymorphic sites in the EDG6 gene, all of which are believed to be novel. The novel EDG6 polymorphic sites identified by the inventors are referred to as PS1-PS23 to designate the order in 5 which they are located in the gene (see Table 3 below). Using the genotypes identified in the Index Repository for PS1-PS23 and the methodology described in the Examples below, the inventors herein also determined the pair of haplotypes for the EDG6 gene present in individual human members of this repository. The human genotypes and haplotypes found in the repository for the EDG6 gene include 10 those shown in Tables 4 and 5, respectively. The polymorphism and haplotype data disclosed herein are useful for validating whether EDG6 is a suitable target for drugs to treat cancer, angiogenesis and inflammation, screening for such drugs and reducing bias in clinical trials of such drugs.

In the context of this disclosure, the following terms shall be defined as follows unless otherwise indicated:

Allele - A particular form of a genetic locus, distinguished from other forms by its particular 15 nucleotide sequence.

Candidate Gene - A gene which is hypothesized to be responsible for a disease, condition, or the response to a treatment, or to be correlated with one of these.

Gene - A segment of DNA that contains all the information for the regulated biosynthesis of an RNA product, including promoters, exons, introns, and other untranslated regions that control 20 expression.

Genotype - An unphased 5' to 3' sequence of nucleotide pair(s) found at one or more polymorphic sites in a locus on a pair of homologous chromosomes in an individual. As used herein, genotype includes a full-genotype and/or a sub-genotype as described below.

Full-genotype - The unphased 5' to 3' sequence of nucleotide pairs found at all polymorphic 25 sites examined herein in a locus on a pair of homologous chromosomes in a single individual.

Sub-genotype - The unphased 5' to 3' sequence of nucleotides seen at a subset of the polymorphic sites examined herein in a locus on a pair of homologous chromosomes in a single individual.

Genotyping - A process for determining a genotype of an individual.

Haplotype - A 5' to 3' sequence of nucleotides found at one or more polymorphic sites in a 30 locus on a single chromosome from a single individual. As used herein, haplotype includes a full-haplotype and/or a sub-haplotype as described below.

Full-haplotype - The 5' to 3' sequence of nucleotides found at all polymorphic sites examined herein in a locus on a single chromosome from a single individual.

Sub-haplotype - The 5' to 3' sequence of nucleotides seen at a subset of the polymorphic sites 35 examined herein in a locus on a single chromosome from a single individual.

Haplotype pair - The two haplotypes found for a locus in a single individual.

Haplotyping - A process for determining one or more haplotypes in an individual and includes

use of family pedigrees, molecular techniques and/or statistical inference.

Haplotype data - Information concerning one or more of the following for a specific gene: a listing of the haplotype pairs in each individual in a population; a listing of the different haplotypes in a population; frequency of each haplotype in that or other populations, and any known associations 5 between one or more haplotypes and a trait.

Isoform - A particular form of a gene, mRNA, cDNA or the protein encoded thereby, distinguished from other forms by its particular sequence and/or structure.

Isogene - One of the isoforms of a gene found in a population. An isogene contains all of the polymorphisms present in the particular isoform of the gene.

10 **Isolated** - As applied to a biological molecule such as RNA, DNA, oligonucleotide, or protein, isolated means the molecule is substantially free of other biological molecules such as nucleic acids, proteins, lipids, carbohydrates, or other material such as cellular debris and growth media. Generally, the term "isolated" is not intended to refer to a complete absence of such material or to absence of water, buffers, or salts, unless they are present in amounts that substantially interfere with the methods 15 of the present invention.

Locus - A location on a chromosome or DNA molecule corresponding to a gene or a physical or phenotypic feature.

20 **Naturally-occurring** - A term used to designate that the object it is applied to, e.g., naturally-occurring polynucleotide or polypeptide, can be isolated from a source in nature and which has not been intentionally modified by man.

Nucleotide pair - The nucleotides found at a polymorphic site on the two copies of a chromosome from an individual.

25 **Phased** - As applied to a sequence of nucleotide pairs for two or more polymorphic sites in a locus, phased means the combination of nucleotides present at those polymorphic sites on a single copy of the locus is known.

Polymorphic site (PS) - A position within a locus at which at least two alternative sequences are found in a population, the most frequent of which has a frequency of no more than 99%.

30 **Polymorphic variant** - A gene, mRNA, cDNA, polypeptide or peptide whose nucleotide or amino acid sequence varies from a reference sequence due to the presence of a polymorphism in the gene.

Polymorphism - The sequence variation observed in an individual at a polymorphic site. Polymorphisms include nucleotide substitutions, insertions, deletions and microsatellites and may, but need not, result in detectable differences in gene expression or protein function.

35 **Polymorphism data** - Information concerning one or more of the following for a specific gene: location of polymorphic sites; sequence variation at those sites; frequency of polymorphisms in one or more populations; the different genotypes and/or haplotypes determined for the gene; frequency of one or more of these genotypes and/or haplotypes in one or more populations; any known association(s) between a trait and a genotype or a haplotype for the gene.

Polymorphism Database – A collection of polymorphism data arranged in a systematic or methodical way and capable of being individually accessed by electronic or other means.

Polynucleotide – A nucleic acid molecule comprised of single-stranded RNA or DNA or comprised of complementary, double-stranded DNA.

5 **Population Group** – A group of individuals sharing a common ethnogeographic origin.

Reference Population – A group of subjects or individuals who are predicted to be representative of the genetic variation found in the general population. Typically, the reference population represents the genetic variation in the population at a certainty level of at least 85%, preferably at least 90%, more preferably at least 95% and even more preferably at least 99%.

10 **Single Nucleotide Polymorphism (SNP)** – Typically, the specific pair of nucleotides observed at a single polymorphic site. In rare cases, three or four nucleotides may be found.

Subject – A human individual whose genotypes or haplotypes or response to treatment or disease state are to be determined.

Treatment - A stimulus administered internally or externally to a subject.

15 **Unphased** – As applied to a sequence of nucleotide pairs for two or more polymorphic sites in a locus, unphased means the combination of nucleotides present at those polymorphic sites on a single copy of the locus is not known.

20 As discussed above, information on the identity of genotypes and haplotypes for the EDG6 gene of any particular individual as well as the frequency of such genotypes and haplotypes in any particular population of individuals is expected to be useful for a variety of drug discovery and development applications. Thus, the invention also provides compositions and methods for detecting the novel EDG6 polymorphisms and haplotypes identified herein.

25 The compositions comprise at least one EDG6 genotyping oligonucleotide. In one embodiment, an EDG6 genotyping oligonucleotide is a probe or primer capable of hybridizing to a target region that is located close to, or that contains, one of the novel polymorphic sites described herein. As used herein, the term “oligonucleotide” refers to a polynucleotide molecule having less than about 100 nucleotides. A preferred oligonucleotide of the invention is 10 to 35 nucleotides long. More preferably, the oligonucleotide is between 15 and 30, and most preferably, between 20 and 25 nucleotides in length. The exact length of the oligonucleotide will depend on many factors that are 30 routinely considered and practiced by the skilled artisan. The oligonucleotide may be comprised of any phosphorylation state of ribonucleotides, deoxyribonucleotides, and acyclic nucleotide derivatives, and other functionally equivalent derivatives. Alternatively, oligonucleotides may have a phosphate-free backbone, which may be comprised of linkages such as carboxymethyl, acetamide, carbamate, polyamide (peptide nucleic acid (PNA)) and the like (Varma, R. in Molecular Biology and 35 Biotechnology, A Comprehensive Desk Reference, Ed. R. Meyers, VCH Publishers, Inc. (1995), pages 617-620). Oligonucleotides of the invention may be prepared by chemical synthesis using any suitable methodology known in the art, or may be derived from a biological sample, for example, by restriction digestion. The oligonucleotides may be labeled, according to any technique known in the art,

including use of radiolabels, fluorescent labels, enzymatic labels, proteins, haptens, antibodies, sequence tags and the like.

Genotyping oligonucleotides of the invention must be capable of specifically hybridizing to a target region of an EDG6 polynucleotide, i.e., an EDG6 isogene. As used herein, specific hybridization means the oligonucleotide forms an anti-parallel double-stranded structure with the target region under certain hybridizing conditions, while failing to form such a structure when incubated with a non-target region or a non-EDG6 polynucleotide under the same hybridizing conditions. Preferably, the oligonucleotide specifically hybridizes to the target region under conventional high stringency conditions. The skilled artisan can readily design and test oligonucleotide probes and primers suitable for detecting polymorphisms in the EDG6 gene using the polymorphism information provided herein in conjunction with the known sequence information for the EDG6 gene and routine techniques.

A nucleic acid molecule such as an oligonucleotide or polynucleotide is said to be a "perfect" or "complete" complement of another nucleic acid molecule if every nucleotide of one of the molecules is complementary to the nucleotide at the corresponding position of the other molecule. A nucleic acid molecule is "substantially complementary" to another molecule if it hybridizes to that molecule with sufficient stability to remain in a duplex form under conventional low-stringency conditions. Conventional hybridization conditions are described, for example, by Sambrook J. et al., in *Molecular Cloning, A Laboratory Manual*, 2nd Edition, Cold Spring Harbor Press, Cold Spring Harbor, NY (1989) and by Haymes, B.D. et al. in *Nucleic Acid Hybridization, A Practical Approach*, IRL Press, Washington, D.C. (1985). While perfectly complementary oligonucleotides are preferred for detecting polymorphisms, departures from complete complementarity are contemplated where such departures do not prevent the molecule from specifically hybridizing to the target region. For example, an oligonucleotide primer may have a non-complementary fragment at its 5' end, with the remainder of the primer being complementary to the target region. Alternatively, non-complementary nucleotides may be interspersed into the oligonucleotide probe or primer as long as the resulting probe or primer is still capable of specifically hybridizing to the target region.

Preferred genotyping oligonucleotides of the invention are allele-specific oligonucleotides. As used herein, the term allele-specific oligonucleotide (ASO) means an oligonucleotide that is able, under sufficiently stringent conditions, to hybridize specifically to one allele of a gene, or other locus, at a target region containing a polymorphic site while not hybridizing to the corresponding region in another allele(s). As understood by the skilled artisan, allele-specificity will depend upon a variety of readily optimized stringency conditions, including salt and formamide concentrations, as well as temperatures for both the hybridization and washing steps. Examples of hybridization and washing conditions typically used for ASO probes are found in Kogan et al., "Genetic Prediction of Hemophilia A" in *PCR Protocols, A Guide to Methods and Applications*, Academic Press, 1990 and Ruaño et al., 87 *Proc. Natl. Acad. Sci. USA* 6296-6300, 1990. Typically, an ASO will be perfectly complementary to one allele while containing a single mismatch for another allele.

Allele-specific oligonucleotides of the invention include ASO probes and ASO primers. ASO probes which usually provide good discrimination between different alleles are those in which a central position of the oligonucleotide probe aligns with the polymorphic site in the target region (e.g., approximately the 7th or 8th position in a 15mer, the 8th or 9th position in a 16mer, and the 10th or 11th position in a 20mer). An ASO primer of the invention has a 3' terminal nucleotide, or preferably a 3' penultimate nucleotide, that is complementary to only one nucleotide of a particular SNP, thereby acting as a primer for polymerase-mediated extension only if the allele containing that nucleotide is present. ASO probes and primers hybridizing to either the coding or noncoding strand are contemplated by the invention.

ASO probes and primers listed below use the appropriate nucleotide symbol (R= G or A, Y= T or C, M= A or C, K= G or T, S= G or C, and W= A or T; WIPO standard ST.25) at the position of the polymorphic site to represent the two alternative allelic variants observed at that polymorphic site.

A preferred ASO probe for detecting EDG6 gene polymorphisms comprises a nucleotide sequence, listed 5' to 3', selected from the group consisting of:

15 GTGTGCTRAGCGCCG (SEQ ID NO:4) and its complement,
 GGCCCATYCCGAGTG (SEQ ID NO:5) and its complement,
 GGGGGTCYTCACAGC (SEQ ID NO:6) and its complement,
 CCAGGGCAGCCCCAG (SEQ ID NO:7) and its complement,
 CCGGGCGYGGGGGGC (SEQ ID NO:8) and its complement,
 20 TGCAGTCRGCACGCT (SEQ ID NO:9) and its complement,
 CGAGAGCAGGGCCAC (SEQ ID NO:10) and its complement,
 CGTCTACRGCTTCAT (SEQ ID NO:11) and its complement,
 TGGCCGCRCTGCTGG (SEQ ID NO:12) and its complement,
 CCTGTGCKCCTTGA (SEQ ID NO:13) and its complement,
 25 AGCGGCCYGCCGCAA (SEQ ID NO:14) and its complement,
 CACTCTTYGGGCTGC (SEQ ID NO:15) and its complement,
 CCGACAGYTCTCTGA (SEQ ID NO:16) and its complement,
 GGCTCCCCCTCGCTC (SEQ ID NO:17) and its complement,
 CTCCAGCRTGCGGAG (SEQ ID NO:18) and its complement,
 30 GTCTTGCRGTGGAT (SEQ ID NO:19) and its complement,
 TCTTCCCRGTGGCCT (SEQ ID NO:20) and its complement,
 CAAATGGRCTTCCA (SEQ ID NO:21) and its complement,
 GATTCTGSGGAAGTC (SEQ ID NO:22) and its complement,
 ATGTTGCRGCCTCTT (SEQ ID NO:23) and its complement,
 35 CTGGTGCRGATGC (SEQ ID NO:24) and its complement,
 GTGCATGYGTGGGGG (SEQ ID NO:25) and its complement, and
 GGCTCAGSGGGGCTG (SEQ ID NO:26) and its complement.

A preferred ASO primer for detecting EDG6 gene polymorphisms comprises a nucleotide sequence, listed 5' to 3', selected from the group consisting of:

40 CCTGCTGTGTGCTRA (SEQ ID NO:27); CTCCACCGGCGCTYA (SEQ ID NO:28);
 AGGGGTGGCCCATYC (SEQ ID NO:29); AGTCCCCACTCGGRA (SEQ ID NO:30);
 AGGGGTGGGGGTCYT (SEQ ID NO:31); GCCCTGGCTGTGARG (SEQ ID NO:32);
 45 TCACAGCCAGGGCRG (SEQ ID NO:33); AACGCGCTGGGGCYG (SEQ ID NO:34);
 GGCTGGCCGGCGYG (SEQ ID NO:35); CCTCCGGCCCCCRC (SEQ ID NO:36);
 GCCACATGCGGTCRC (SEQ ID NO:37); AGACCCAGCGTGYG (SEQ ID NO:38);
 GGTGGCCGAGAGCRG (SEQ ID NO:39); GTCTTGGTGGCCYG (SEQ ID NO:40);
 CAGCCGCGTCTACRG (SEQ ID NO:41); AGGCCGATGAAGCYG (SEQ ID NO:42);

GGCTGCTGGCCGCRC (SEQ ID NO:43); GCATCCCCAGCAGYG (SEQ ID NO:44);
 GAACTGCCTGTGCKC (SEQ ID NO:45); CAGCGGTCAAAGGMG (SEQ ID NO:46);
 ACGCCCAGCGGCCYG (SEQ ID NO:47); CGGGCCTTGCAGCRG (SEQ ID NO:48);
 5 GGGGCCCACTCTTYG (SEQ ID NO:49); CCAGCAGCAGCCCRA (SEQ ID NO:50);
 CCACCACCGACAGYT (SEQ ID NO:51); TTGGCCTCAGAGARC (SEQ ID NO:52);
 TTTCGCGGCTCCCRC (SEQ ID NO:53); AAAGCTGAGCGAGMG (SEQ ID NO:54);
 CAGCATCTCCAGCRT (SEQ ID NO:55); CAGATGCTCCGCAYG (SEQ ID NO:56);
 GTTGCAGTCTGCRT (SEQ ID NO:57); TGCACCACATCCACAYG (SEQ ID NO:58);
 CCATGGTCTTCCCRG (SEQ ID NO:59); CCCGAGAGGCCACYG (SEQ ID NO:60);
 10 TGACGCCAAATGGRC (SEQ ID NO:61); TGACCATGGGAAGYC (SEQ ID NO:62);
 CTGTGTGATTCTGSG (SEQ ID NO:63); GGCCGGGACTTCSC (SEQ ID NO:64);
 TACGTGATGTTGCRG (SEQ ID NO:65); GGGAAATAAGAGGCY (SEQ ID NO:66);
 TATTCCCTGGTGCRT (SEQ ID NO:67); CCCCACGCATGCAYG (SEQ ID NO:68);
 TGGTGCCTGCATGYG (SEQ ID NO:69); CCACGGCCCCCACRC (SEQ ID NO:70);
 15 GGCGTGGCTCAGSG (SEQ ID NO:71); and GATCCACAGCCCCSC (SEQ ID NO:72).

Other genotyping oligonucleotides of the invention hybridize to a target region located one to several nucleotides downstream of one of the novel polymorphic sites identified herein. Such 20 oligonucleotides are useful in polymerase-mediated primer extension methods for detecting one of the novel polymorphisms described herein and therefore such genotyping oligonucleotides are referred to herein as "primer-extension oligonucleotides". In a preferred embodiment, the 3'-terminus of a primer-extension oligonucleotide is a deoxynucleotide complementary to the nucleotide located immediately adjacent to the polymorphic site.

25 A particularly preferred oligonucleotide primer for detecting EDG6 gene polymorphisms by primer extension terminates in a nucleotide sequence, listed 5' to 3', selected from the group consisting of:

30 GCTGTGTGCT (SEQ ID NO:73); CACCGGCGCT (SEQ ID NO:74);
 GGTGGCCCAT (SEQ ID NO:75); CCCCCACTCGG (SEQ ID NO:76);
 GGTGGGGGTC (SEQ ID NO:77); CTGGCTGTGA (SEQ ID NO:78);
 CAGCCAGGGC (SEQ ID NO:79); GCGCTGGGGC (SEQ ID NO:80);
 TGGCCGGGCG (SEQ ID NO:81); CGGGCCCCCC (SEQ ID NO:82);
 ACATGCGGTC (SEQ ID NO:83); CCCAGCGTCG (SEQ ID NO:84);
 35 GGCCGAGAGC (SEQ ID NO:85); TTGGTGGGCC (SEQ ID NO:86);
 CCGCGTCTAC (SEQ ID NO:87); CCGATGAAGC (SEQ ID NO:88);
 TGCTGGCCGC (SEQ ID NO:89); TCCCCAGCAG (SEQ ID NO:90);
 CTGCCTGTGC (SEQ ID NO:91); CGGTCAAAGG (SEQ ID NO:92);
 40 CCCAGCGGCC (SEQ ID NO:93); GCCTTGCAGC (SEQ ID NO:94);
 GCCCACTCTT (SEQ ID NO:95); GCAGCAGCCC (SEQ ID NO:96);
 CCACCGACAG (SEQ ID NO:97); GCCTCAGAGA (SEQ ID NO:98);
 CGCGGCTCCC (SEQ ID NO:99); GCTGAGCGAG (SEQ ID NO:100);
 CATCTCCAGC (SEQ ID NO:101); ATGCTCCGCA (SEQ ID NO:102);
 45 GCAGTCTTGC (SEQ ID NO:103); ACCATCCACA (SEQ ID NO:104);
 TGGTCTTCCC (SEQ ID NO:105); GAGAGGCCAC (SEQ ID NO:106);
 CGCCAAATGG (SEQ ID NO:107); CCATGGGAAG (SEQ ID NO:108);
 TGTGATTCTG (SEQ ID NO:109); CGGGACTTCC (SEQ ID NO:110);
 GTGATGTTGC (SEQ ID NO:111); AATAAGAGGC (SEQ ID NO:112);
 50 TCCCTGGTGC (SEQ ID NO:113); CACGCATGCA (SEQ ID NO:114);
 TGCGTGCATG (SEQ ID NO:115); CGGGCCCCCAC (SEQ ID NO:116);
 CGTGGCTCAG (SEQ ID NO:117); and CCACAGCCCC (SEQ ID NO:118).

5 In some embodiments, a composition contains two or more differently labeled genotyping oligonucleotides for simultaneously probing the identity of nucleotides at two or more polymorphic sites. It is also contemplated that primer compositions may contain two or more sets of allele-specific primer pairs to allow simultaneous targeting and amplification of two or more regions containing a polymorphic site.

10 EDG6 genotyping oligonucleotides of the invention may also be immobilized on or synthesized on a solid surface such as a microchip, bead, or glass slide (see, e.g., WO 98/20020 and WO 98/20019). Such immobilized genotyping oligonucleotides may be used in a variety of polymorphism detection assays, including but not limited to probe hybridization and polymerase extension assays. Immobilized EDG6 genotyping oligonucleotides of the invention may comprise an ordered array of oligonucleotides designed to rapidly screen a DNA sample for polymorphisms in multiple genes at the same time.

15 In another embodiment, the invention provides a kit comprising at least two genotyping oligonucleotides packaged in separate containers. The kit may also contain other components such as hybridization buffer (where the oligonucleotides are to be used as a probe) packaged in a separate container. Alternatively, where the oligonucleotides are to be used to amplify a target region, the kit may contain, packaged in separate containers, a polymerase and a reaction buffer optimized for primer extension mediated by the polymerase, such as PCR.

20 The above described oligonucleotide compositions and kits are useful in methods for genotyping and/or haplotyping the EDG6 gene in an individual. As used herein, the terms "EDG6 genotype" and "EDG6 haplotype" mean the genotype or haplotype contains the nucleotide pair or nucleotide, respectively, that is present at one or more of the novel polymorphic sites described herein and may optionally also include the nucleotide pair or nucleotide present at one or more additional 25 polymorphic sites in the EDG6 gene. The additional polymorphic sites may be currently known polymorphic sites or sites that are subsequently discovered.

30 One embodiment of the genotyping method involves isolating from the individual a nucleic acid sample comprising the two copies of the EDG6 gene, or a fragment thereof, that are present in the individual, and determining the identity of the nucleotide pair at one or more polymorphic sites selected from the group consisting of PS1, PS2, PS3, PS4, PS5, PS6, PS7, PS8, PS9, PS10, PS11, PS12, PS13, PS14, PS15, PS16, PS17, PS18, PS19, PS20, PS21, PS22 and PS23 in the two copies to 35 assign an EDG6 genotype to the individual. As will be readily understood by the skilled artisan, the two "copies" of a gene in an individual may be the same allele or may be different alleles. In a particularly preferred embodiment, the genotyping method comprises determining the identity of the nucleotide pair at each of PS1-PS23.

Typically, the nucleic acid sample is isolated from a biological sample taken from the individual, such as a blood sample or tissue sample. Suitable tissue samples include whole blood, semen, saliva, tears, urine, fecal material, sweat, buccal, skin and hair. The nucleic acid sample may

be comprised of genomic DNA, mRNA, or cDNA and, in the latter two cases, the biological sample must be obtained from a tissue in which the EDG6 gene is expressed. Furthermore it will be understood by the skilled artisan that mRNA or cDNA preparations would not be used to detect polymorphisms located in introns or in 5' and 3' untranslated regions. If an EDG6 gene fragment is 5 isolated, it must contain the polymorphic site(s) to be genotyped.

One embodiment of the haplotyping method comprises isolating from the individual a nucleic acid sample containing only one of the two copies of the EDG6 gene, or a fragment thereof, that is present in the individual and determining in that copy the identity of the nucleotide at one or more polymorphic sites selected from the group consisting of PS1, PS2, PS3, PS4, PS5, PS6, PS7, PS8, PS9, 10 PS10, PS11, PS12, PS13, PS14, PS15, PS16, PS17, PS18, PS19, PS20, PS21, PS22 and PS23 in that copy to assign an EDG6 haplotype to the individual. The nucleic acid may be isolated using any method capable of separating the two copies of the EDG6 gene or fragment such as one of the methods described above for preparing EDG6 isogenes, with targeted *in vivo* cloning being the preferred approach. As will be readily appreciated by those skilled in the art, any individual clone will only 15 provide haplotype information on one of the two EDG6 gene copies present in an individual. If haplotype information is desired for the individual's other copy, additional EDG6 clones will need to be examined. Typically, at least five clones should be examined to have more than a 90% probability of haplotyping both copies of the EDG6 gene in an individual. In a particularly preferred embodiment, the nucleotide at each of PS1-PS23 is identified.

20 In another embodiment, the haplotyping method comprises determining whether an individual has one or more of the EDG6 haplotypes shown in Table 5. This can be accomplished by identifying, for one or both copies of the individual's EDG6 gene, the phased sequence of nucleotides present at each of PS1-PS23. The present invention also contemplates that typically only a subset of PS1-PS23 will need to be directly examined to assign to an individual one or more of the haplotypes shown in 25 Table 5. This is because at least one polymorphic site in a gene is frequently in strong linkage disequilibrium with one or more other polymorphic sites in that gene (Drysdale, CM et al. 2000 *PNAS* 97:10483-10488; Rieder MJ et al. 1999 *Nature Genetics* 22:59-62). Two sites are said to be in linkage disequilibrium if the presence of a particular variant at one site enhances the predictability of another variant at the second site (Stephens, JC 1999, *Mol. Diag.* 4:309-317). Techniques for determining 30 whether any two polymorphic sites are in linkage disequilibrium are well-known in the art (Weir B.S. 1996 *Genetic Data Analysis II*, Sinauer Associates, Inc. Publishers, Sunderland, MA).

In a preferred embodiment, an EDG6 haplotype pair is determined for an individual by identifying the phased sequence of nucleotides at one or more polymorphic sites selected from the group consisting of PS1, PS2, PS3, PS4, PS5, PS6, PS7, PS8, PS9, PS10, PS11, PS12, PS13, PS14, 35 PS15, PS16, PS17, PS18, PS19, PS20, PS21, PS22 and PS23 in each copy of the EDG6 gene that is present in the individual. In a particularly preferred embodiment, the haplotyping method comprises identifying the phased sequence of nucleotides at each of PS1-PS23 in each copy of the EDG6 gene. When haplotyping both copies of the gene, the identifying step is preferably performed with each copy

of the gene being placed in separate containers. However, it is also envisioned that if the two copies are labeled with different tags, or are otherwise separately distinguishable or identifiable, it could be possible in some cases to perform the method in the same container. For example, if first and second copies of the gene are labeled with different first and second fluorescent dyes, respectively, and an 5 allele-specific oligonucleotide labeled with yet a third different fluorescent dye is used to assay the polymorphic site(s), then detecting a combination of the first and third dyes would identify the polymorphism in the first gene copy while detecting a combination of the second and third dyes would identify the polymorphism in the second gene copy.

In both the genotyping and haplotyping methods, the identity of a nucleotide (or nucleotide 10 pair) at a polymorphic site(s) may be determined by amplifying a target region(s) containing the polymorphic site(s) directly from one or both copies of the EDG6 gene, or a fragment thereof, and the sequence of the amplified region(s) determined by conventional methods. It will be readily appreciated by the skilled artisan that only one nucleotide will be detected at a polymorphic site in individuals who are homozygous at that site, while two different nucleotides will be detected if the 15 individual is heterozygous for that site. The polymorphism may be identified directly, known as positive-type identification, or by inference, referred to as negative-type identification. For example, where a SNP is known to be guanine and cytosine in a reference population, a site may be positively determined to be either guanine or cytosine for an individual homozygous at that site, or both guanine and cytosine, if the individual is heterozygous at that site. Alternatively, the site may be negatively 20 determined to be not guanine (and thus cytosine/cytosine) or not cytosine (and thus guanine/guanine).

The target region(s) may be amplified using any oligonucleotide-directed amplification method, including but not limited to polymerase chain reaction (PCR) (U.S. Patent No. 4,965,188), 25 ligase chain reaction (LCR) (Barany et al., *Proc. Natl. Acad. Sci. USA* 88:189-193, 1991; WO90/01069), and oligonucleotide ligation assay (OLA) (Landegren et al., *Science* 241:1077-1080, 1988).

Other known nucleic acid amplification procedures may be used to amplify the target region including transcription-based amplification systems (U.S. Patent No. 5,130,238; EP 329,822; U.S. Patent No. 5,169,766, WO89/06700) and isothermal methods (Walker et al., *Proc. Natl. Acad. Sci. USA* 89:392-396, 1992).

30 A polymorphism in the target region may also be assayed before or after amplification using one of several hybridization-based methods known in the art. Typically, allele-specific oligonucleotides are utilized in performing such methods. The allele-specific oligonucleotides may be used as differently labeled probe pairs, with one member of the pair showing a perfect match to one variant of a target sequence and the other member showing a perfect match to a different variant. In 35 some embodiments, more than one polymorphic site may be detected at once using a set of allele-specific oligonucleotides or oligonucleotide pairs. Preferably, the members of the set have melting temperatures within 5°C, and more preferably within 2°C, of each other when hybridizing to each of the polymorphic sites being detected.

Hybridization of an allele-specific oligonucleotide to a target polynucleotide may be performed with both entities in solution, or such hybridization may be performed when either the oligonucleotide or the target polynucleotide is covalently or noncovalently affixed to a solid support. Attachment may be mediated, for example, by antibody-antigen interactions, poly-L-Lys, streptavidin or avidin-biotin, 5 salt bridges, hydrophobic interactions, chemical linkages, UV cross-linking baking, etc. Allele-specific oligonucleotides may be synthesized directly on the solid support or attached to the solid support subsequent to synthesis. Solid-supports suitable for use in detection methods of the invention include substrates made of silicon, glass, plastic, paper and the like, which may be formed, for example, into wells (as in 96-well plates), slides, sheets, membranes, fibers, chips, dishes, and beads. 10 The solid support may be treated, coated or derivatized to facilitate the immobilization of the allele-specific oligonucleotide or target nucleic acid.

The genotype or haplotype for the EDG6 gene of an individual may also be determined by hybridization of a nucleic acid sample containing one or both copies of the gene, or fragment(s) thereof, to nucleic acid arrays and subarrays such as described in WO 95/11995. The arrays would 15 contain a battery of allele-specific oligonucleotides representing each of the polymorphic sites to be included in the genotype or haplotype.

The identity of polymorphisms may also be determined using a mismatch detection technique, including but not limited to the RNase protection method using riboprobes (Winter et al., *Proc. Natl. Acad. Sci. USA* 82:7575, 1985; Meyers et al., *Science* 230:1242, 1985) and proteins which recognize 20 nucleotide mismatches, such as the *E. coli* mutS protein (Modrich, P. *Ann. Rev. Genet.* 25:229-253, 1991). Alternatively, variant alleles can be identified by single strand conformation polymorphism (SSCP) analysis (Orita et al., *Genomics* 5:874-879, 1989; Humphries et al., in *Molecular Diagnosis of Genetic Diseases*, R. Elles, ed., pp. 321-340, 1996) or denaturing gradient gel electrophoresis (DGGE) (Wartell et al., *Nucl. Acids Res.* 18:2699-2706, 1990; Sheffield et al., *Proc. Natl. Acad. Sci. USA* 25:86:232-236, 1989).

A polymerase-mediated primer extension method may also be used to identify the polymorphism(s). Several such methods have been described in the patent and scientific literature and include the "Genetic Bit Analysis" method (WO92/15712) and the ligase/polymerase mediated genetic bit analysis (U.S. Patent 5,679,524. Related methods are disclosed in WO91/02087, WO90/09455, 30 WO95/17676, U.S. Patent Nos. 5,302,509, and 5,945,283. Extended primers containing a polymorphism may be detected by mass spectrometry as described in U.S. Patent No. 5,605,798. Another primer extension method is allele-specific PCR (Ruaño et al., *Nucl. Acids Res.* 17:8392, 1989; Ruaño et al., *Nucl. Acids Res.* 19, 6877-6882, 1991; WO 93/22456; Turki et al., *J. Clin. Invest.* 95:1635-1641, 1995). In addition, multiple polymorphic sites may be investigated by simultaneously 35 amplifying multiple regions of the nucleic acid using sets of allele-specific primers as described in Wallace et al. (WO89/10414).

In addition, the identity of the allele(s) present at any of the novel polymorphic sites described herein may be indirectly determined by genotyping another polymorphic site that is in linkage

disequilibrium with the polymorphic site that is of interest. Polymorphic sites in linkage disequilibrium with the presently disclosed polymorphic sites may be located in regions of the gene or in other genomic regions not examined herein. Genotyping of a polymorphic site in linkage disequilibrium with the novel polymorphic sites described herein may be performed by, but is not limited to, any of the above-mentioned methods for detecting the identity of the allele at a polymorphic site.

In another aspect of the invention, an individual's EDG6 haplotype pair is predicted from its EDG6 genotype using information on haplotype pairs known to exist in a reference population. In its broadest embodiment, the haplotyping prediction method comprises identifying an EDG6 genotype for the individual at two or more EDG6 polymorphic sites described herein, enumerating all possible haplotype pairs which are consistent with the genotype, accessing data containing EDG6 haplotype pairs identified in a reference population, and assigning a haplotype pair to the individual that is consistent with the data. In one embodiment, the reference haplotype pairs include the EDG6 haplotype pairs shown in Table 4.

Generally, the reference population should be composed of randomly-selected individuals representing the major ethnogeographic groups of the world. A preferred reference population for use in the methods of the present invention comprises an approximately equal number of individuals from Caucasian, African-descent, Asian and Hispanic-Latino population groups with the minimum number of each group being chosen based on how rare a haplotype one wants to be guaranteed to see. For example, if one wants to have a q% chance of not missing a haplotype that exists in the population at a p% frequency of occurring in the reference population, the number of individuals (n) who must be sampled is given by $2n = \log(1-q)/\log(1-p)$ where p and q are expressed as fractions. A preferred reference population allows the detection of any haplotype whose frequency is at least 10% with about 99% certainty and comprises about 20 unrelated individuals from each of the four population groups named above. A particularly preferred reference population includes a 3-generation family representing one or more of the four population groups to serve as controls for checking quality of haplotyping procedures.

In a preferred embodiment, the haplotype frequency data for each ethnogeographic group is examined to determine whether it is consistent with Hardy-Weinberg equilibrium. Hardy-Weinberg equilibrium (D.L. Hartl et al., *Principles of Population Genetics*, Sinauer Associates (Sunderland, MA), 3rd Ed., 1997) postulates that the frequency of finding the haplotype pair H_1 / H_2 is equal to $p_{H-W}(H_1 / H_2) = 2p(H_1)p(H_2)$ if $H_1 \neq H_2$ and $p_{H-W}(H_1 / H_2) = p(H_1)p(H_2)$ if $H_1 = H_2$. A statistically significant difference between the observed and expected haplotype frequencies could be due to one or more factors including significant inbreeding in the population group, strong selective pressure on the gene, sampling bias, and/or errors in the genotyping process. If large deviations from Hardy-Weinberg equilibrium are observed in an ethnogeographic group, the number of individuals in that group can be increased to see if the deviation is due to a sampling bias. If a larger sample size

does not reduce the difference between observed and expected haplotype pair frequencies, then one may wish to consider haplotyping the individual using a direct haplotyping method such as, for example, CLASPER System™ technology (U.S. Patent No. 5,866,404), single molecule dilution, or allele-specific long-range PCR (Michalotos-Beloin et al., *Nucleic Acids Res.* 24:4841-4843, 1996).

5 In one embodiment of this method for predicting an EDG6 haplotype pair for an individual, the assigning step involves performing the following analysis. First, each of the possible haplotype pairs is compared to the haplotype pairs in the reference population. Generally, only one of the haplotype pairs in the reference population matches a possible haplotype pair and that pair is assigned to the individual. Occasionally, only one haplotype represented in the reference haplotype pairs is
10 consistent with a possible haplotype pair for an individual, and in such cases the individual is assigned a haplotype pair containing this known haplotype and a new haplotype derived by subtracting the known haplotype from the possible haplotype pair. Alternatively, the haplotype pair in an individual may be predicted from the individual's genotype for that gene using reported methods (e.g., Clark et al. 1990 *Mol Bio Evol* 7:111-22) or through a commercial haplotyping service such as offered by
15 Genaissance Pharmaceuticals, Inc. (New Haven, CT). In rare cases, either no haplotypes in the reference population are consistent with the possible haplotype pairs, or alternatively, multiple reference haplotype pairs are consistent with the possible haplotype pairs. In such cases, the individual is preferably haplotyped using a direct molecular haplotyping method such as, for example, CLASPER System™ technology (U.S. Patent No. 5,866,404), SMD, or allele-specific long-range PCR
20 (Michalotos-Beloin et al., *supra*). A preferred process for predicting EDG6 haplotype pairs from EDG6 genotypes is described in U.S. Provisional Application Serial No. 60/198,340 and the corresponding International Application, PCT/US01/12831.

The invention also provides a method for determining the frequency of an EDG6 genotype, haplotype, or haplotype pair in a population. The method comprises, for each member of the
25 population, determining the genotype or the haplotype pair for the novel EDG6 polymorphic sites described herein, and calculating the frequency any particular genotype, haplotype, or haplotype pair is found in the population. The population may be a reference population, a family population, a same sex population, a population group, or a trait population (e.g., a group of individuals exhibiting a trait of interest such as a medical condition or response to a therapeutic treatment).

30 In another aspect of the invention, frequency data for EDG6 genotypes, haplotypes, and/or haplotype pairs are determined in a reference population and used in a method for identifying an association between a trait and an EDG6 genotype, haplotype, or haplotype pair. The trait may be any detectable phenotype, including but not limited to susceptibility to a disease or response to a treatment. The method involves obtaining data on the frequency of the genotype(s), haplotype(s), or haplotype
35 pair(s) of interest in a reference population as well as in a population exhibiting the trait. Frequency data for one or both of the reference and trait populations may be obtained by genotyping or haplotyping each individual in the populations using one of the methods described above. The haplotypes for the trait population may be determined directly or, alternatively, by the predictive

genotype to haplotype approach described above. In another embodiment, the frequency data for the reference and/or trait populations is obtained by accessing previously determined frequency data, which may be in written or electronic form. For example, the frequency data may be present in a database that is accessible by a computer. Once the frequency data is obtained, the frequencies of the 5 genotype(s), haplotype(s), or haplotype pair(s) of interest in the reference and trait populations are compared. In a preferred embodiment, the frequencies of all genotypes, haplotypes, and/or haplotype pairs observed in the populations are compared. If a particular EDG6 genotype, haplotype, or haplotype pair is more frequent in the trait population than in the reference population at a statistically significant amount, then the trait is predicted to be associated with that EDG6 genotype, haplotype or 10 haplotype pair. Preferably, the EDG6 genotype, haplotype, or haplotype pair being compared in the trait and reference populations is selected from the full-genotypes and full-haplotypes shown in Tables 4 and 5, or from sub-genotypes and sub-haplotypes derived from these genotypes and haplotypes.

In a preferred embodiment of the method, the trait of interest is a clinical response exhibited by a patient to some therapeutic treatment, for example, response to a drug targeting EDG6 or response 15 to a therapeutic treatment for a medical condition. As used herein, “medical condition” includes but is not limited to any condition or disease manifested as one or more physical and/or psychological symptoms for which treatment is desirable, and includes previously and newly identified diseases and other disorders. As used herein the term “clinical response” means any or all of the following: a quantitative measure of the response, no response, and adverse response (i.e., side effects).

20 In order to deduce a correlation between clinical response to a treatment and an EDG6 genotype, haplotype, or haplotype pair, it is necessary to obtain data on the clinical responses exhibited by a population of individuals who received the treatment, hereinafter the “clinical population”. This clinical data may be obtained by analyzing the results of a clinical trial that has already been run and/or the clinical data may be obtained by designing and carrying out one or more new clinical trials. As 25 used herein, the term “clinical trial” means any research study designed to collect clinical data on responses to a particular treatment, and includes but is not limited to phase I, phase II and phase III clinical trials. Standard methods are used to define the patient population and to enroll subjects.

It is preferred that the individuals included in the clinical population have been graded for the 30 existence of the medical condition of interest. This is important in cases where the symptom(s) being presented by the patients can be caused by more than one underlying condition, and where treatment of the underlying conditions are not the same. An example of this would be where patients experience breathing difficulties that are due to either asthma or respiratory infections. If both sets were treated with an asthma medication, there would be a spurious group of apparent non-responders that did not actually have asthma. These people would affect the ability to detect any correlation between 35 haplotype and treatment outcome. This grading of potential patients could employ a standard physical exam or one or more lab tests. Alternatively, grading of patients could use haplotyping for situations where there is a strong correlation between haplotype pair and disease susceptibility or severity.

The therapeutic treatment of interest is administered to each individual in the trial population

and each individual's response to the treatment is measured using one or more predetermined criteria. It is contemplated that in many cases, the trial population will exhibit a range of responses and that the investigator will choose the number of responder groups (e.g., low, medium, high) made up by the various responses. In addition, the EDG6 gene for each individual in the trial population is genotyped and/or haplotyped, which may be done before or after administering the treatment.

After both the clinical and polymorphism data have been obtained, correlations between individual response and EDG6 genotype or haplotype content are created. Correlations may be produced in several ways. In one method, individuals are grouped by their EDG6 genotype or haplotype (or haplotype pair) (also referred to as a polymorphism group), and then the averages and standard deviations of clinical responses exhibited by the members of each polymorphism group are calculated.

These results are then analyzed to determine if any observed variation in clinical response between polymorphism groups is statistically significant. Statistical analysis methods which may be used are described in L.D. Fisher and G. vanBelle, "Biostatistics: A Methodology for the Health Sciences", Wiley-Interscience (New York) 1993. This analysis may also include a regression calculation of which polymorphic sites in the EDG6 gene give the most significant contribution to the differences in phenotype. One regression model useful in the invention is described in PCT Application Serial No. PCT/US00/17540, entitled "Methods for Obtaining and Using Haplotype Data".

A second method for finding correlations between EDG6 haplotype content and clinical responses uses predictive models based on error-minimizing optimization algorithms. One of many possible optimization algorithms is a genetic algorithm (R. Judson, "Genetic Algorithms and Their Uses in Chemistry" in *Reviews in Computational Chemistry*, Vol. 10, pp. 1-73, K. B. Lipkowitz and D. B. Boyd, eds. (VCH Publishers, New York, 1997). Simulated annealing (Press et al., "Numerical Recipes in C: The Art of Scientific Computing", Cambridge University Press (Cambridge) 1992, Ch. 10), neural networks (E. Rich and K. Knight, "Artificial Intelligence", 2nd Edition (McGraw-Hill, New York, 1991, Ch. 18), standard gradient descent methods (Press et al., *supra*, Ch. 10), or other global or local optimization approaches (see discussion in Judson, *supra*) could also be used. Preferably, the correlation is found using a genetic algorithm approach as described in PCT Application Serial No. PCT/US00/17540.

Correlations may also be analyzed using analysis of variation (ANOVA) techniques to determine how much of the variation in the clinical data is explained by different subsets of the polymorphic sites in the EDG6 gene. As described in PCT Application Serial No. PCT/US00/17540, ANOVA is used to test hypotheses about whether a response variable is caused by or correlated with one or more traits or variables that can be measured (Fisher and vanBelle, *supra*, Ch. 10).

From the analyses described above, a mathematical model may be readily constructed by the skilled artisan that predicts clinical response as a function of EDG6 genotype or haplotype content. Preferably, the model is validated in one or more follow-up clinical trials designed to test the model.

The identification of an association between a clinical response and a genotype or haplotype (or haplotype pair) for the EDG6 gene may be the basis for designing a diagnostic method to determine those individuals who will or will not respond to the treatment, or alternatively, will respond at a lower level and thus may require more treatment, i.e., a greater dose of a drug. The diagnostic method may 5 take one of several forms: for example, a direct DNA test (i.e., genotyping or haplotyping one or more of the polymorphic sites in the EDG6 gene), a serological test, or a physical exam measurement. The only requirement is that there be a good correlation between the diagnostic test results and the underlying EDG6 genotype or haplotype that is in turn correlated with the clinical response. In a preferred embodiment, this diagnostic method uses the predictive haplotyping method described 10 above.

In another embodiment, the invention provides an isolated polynucleotide comprising a polymorphic variant of the EDG6 gene or a fragment of the gene which contains at least one of the novel polymorphic sites described herein. The nucleotide sequence of a variant EDG6 gene is identical to the reference genomic sequence for those portions of the gene examined, as described in 15 the Examples below, except that it comprises a different nucleotide at one or more of the novel polymorphic sites PS1, PS2, PS3, PS4, PS5, PS6, PS7, PS8, PS9, PS10, PS11, PS12, PS13, PS14, PS15, PS16, PS17, PS18, PS19, PS20, PS21, PS22 and PS23. Similarly, the nucleotide sequence of a variant fragment of the EDG6 gene is identical to the corresponding portion of the reference sequence except for having a different nucleotide at one or more of the novel polymorphic sites described herein. 20 Thus, the invention specifically does not include polynucleotides comprising a nucleotide sequence identical to the reference sequence of the EDG6 gene, which is defined by haplotype 5, (or other reported EDG6 sequences) or to portions of the reference sequence (or other reported EDG6 sequences), except for genotyping oligonucleotides as described above.

The location of a polymorphism in a variant gene or fragment is identified by aligning its 25 sequence against SEQ ID NO:1. The polymorphism is selected from the group consisting of adenine at PS1, thymine at PS2, thymine at PS3, guanine at PS4, thymine at PS5, adenine at PS6, adenine at PS7, adenine at PS8, adenine at PS9, thymine at PS10, thymine at PS11, thymine at PS12, thymine at PS13, thymine at PS14, adenine at PS15, adenine at PS16, adenine at PS17, adenine at PS18, cytosine at PS19, adenine at PS20, adenine at PS21, thymine at PS22 and cytosine at PS23. In a preferred 30 embodiment, the polymorphic variant comprises a naturally-occurring isogene of the EDG6 gene which is defined by any one of haplotypes 1- 4 and 6 - 24 shown in Table 5 below.

Polymorphic variants of the invention may be prepared by isolating a clone containing the 35 EDG6 gene from a human genomic library. The clone may be sequenced to determine the identity of the nucleotides at the novel polymorphic sites described herein. Any particular variant claimed herein could be prepared from this clone by performing *in vitro* mutagenesis using procedures well-known in the art.

EDG6 isogenes may be isolated using any method that allows separation of the two "copies" of the EDG6 gene present in an individual, which, as readily understood by the skilled artisan, may be

the same allele or different alleles. Separation methods include targeted *in vivo* cloning (TIVC) in yeast as described in WO 98/01573, U.S. Patent No. 5,866,404, and U.S. Patent No. 5,972,614.

Another method, which is described in U.S. Patent No. 5,972,614, uses an allele specific oligonucleotide in combination with primer extension and exonuclease degradation to generate hemizygous DNA targets. Yet other methods are single molecule dilution (SMD) as described in 5 Ruaño et al., *Proc. Natl. Acad. Sci.* 87:6296-6300, 1990; and allele specific PCR (Ruaño et al., 1989, *supra*; Ruaño et al., 1991, *supra*; Michalatos-Beloin et al., *supra*).

The invention also provides EDG6 genome anthologies, which are collections of EDG6 isogenes found in a given population. The population may be any group of at least two individuals, 10 including but not limited to a reference population, a population group, a family population, a clinical population, and a same sex population. An EDG6 genome anthology may comprise individual EDG6 isogenes stored in separate containers such as microtest tubes, separate wells of a microtitre plate and the like. Alternatively, two or more groups of the EDG6 isogenes in the anthology may be stored in 15 separate containers. Individual isogenes or groups of isogenes in a genome anthology may be stored in any convenient and stable form, including but not limited to in buffered solutions, as DNA precipitates, freeze-dried preparations and the like. A preferred EDG6 genome anthology of the 20 invention comprises a set of isogenes defined by the haplotypes shown in Table 5 below.

An isolated polynucleotide containing a polymorphic variant nucleotide sequence of the 25 invention may be operably linked to one or more expression regulatory elements in a recombinant expression vector capable of being propagated and expressing the encoded EDG6 protein in a prokaryotic or a eukaryotic host cell. Examples of expression regulatory elements which may be used include, but are not limited to, the lac system, operator and promoter regions of phage lambda, yeast 30 promoters, and promoters derived from vaccinia virus, adenovirus, retroviruses, or SV40. Other regulatory elements include, but are not limited to, appropriate leader sequences, termination codons, polyadenylation signals, and other sequences required for the appropriate transcription and subsequent 35 translation of the nucleic acid sequence in a given host cell. Of course, the correct combinations of expression regulatory elements will depend on the host system used. In addition, it is understood that the expression vector contains any additional elements necessary for its transfer to and subsequent replication in the host cell. Examples of such elements include, but are not limited to, origins of 40 replication and selectable markers. Such expression vectors are commercially available or are readily constructed using methods known to those in the art (e.g., F. Ausubel et al., 1987, in "Current Protocols in Molecular Biology", John Wiley and Sons, New York, New York). Host cells which may be used to express the variant EDG6 sequences of the invention include, but are not limited to, eukaryotic and mammalian cells, such as animal, plant, insect and yeast cells, and prokaryotic cells, 45 such as *E. coli*, or algal cells as known in the art. The recombinant expression vector may be introduced into the host cell using any method known to those in the art including, but not limited to, microinjection, electroporation, particle bombardment, transduction, and transfection using DEAE-dextran, lipofection, or calcium phosphate (see e.g., Sambrook et al. (1989) in "Molecular Cloning. A

Laboratory Manual", Cold Spring Harbor Press, Plainview, New York). In a preferred aspect, eukaryotic expression vectors that function in eukaryotic cells, and preferably mammalian cells, are used. Non-limiting examples of such vectors include vaccinia virus vectors, adenovirus vectors, herpes virus vectors, and baculovirus transfer vectors. Preferred eukaryotic cell lines include COS 5 cells, CHO cells, HeLa cells, NIH/3T3 cells, and embryonic stem cells (Thomson, J. A. et al., 1998 *Science* 282:1145-1147). Particularly preferred host cells are mammalian cells.

As will be readily recognized by the skilled artisan, expression of polymorphic variants of the EDG6 gene will produce EDG6 mRNAs varying from each other at any polymorphic site retained in the spliced and processed mRNA molecules. These mRNAs can be used for the preparation of an 10 EDG6 cDNA comprising a nucleotide sequence which is a polymorphic variant of the EDG6 reference coding sequence shown in Figure 2. Thus, the invention also provides EDG6 mRNAs and corresponding cDNAs which comprise a nucleotide sequence that is identical to SEQ ID NO:2 (Fig. 2), or its corresponding RNA sequence, except for having one or more polymorphisms selected from the group consisting of thymine at a position corresponding to nucleotide 114, adenine at a position 15 corresponding to nucleotide 231, adenine at a position corresponding to nucleotide 463, adenine at a position corresponding to nucleotide 490, adenine at a position corresponding to nucleotide 522, thymine at a position corresponding to nucleotide 565, thymine at a position corresponding to nucleotide 727, thymine at a position corresponding to nucleotide 804, thymine at a position corresponding to nucleotide 1059, thymine at a position corresponding to nucleotide 1094 and adenine 20 at a position corresponding to nucleotide 1141. A particularly preferred polymorphic cDNA variant comprises the coding sequence of an EDG6 isogene defined by haplotypes 3c, 7c-12c, 19c-22c, and 24c. Fragments of these variant mRNAs and cDNAs are included in the scope of the invention, provided they contain the novel polymorphisms described herein. The invention specifically excludes 25 polynucleotides identical to previously identified and characterized EDG6 cDNAs and fragments thereof. Polynucleotides comprising a variant RNA or DNA sequence may be isolated from a biological sample using well-known molecular biological procedures or may be chemically synthesized.

As used herein, a polymorphic variant of an EDG6 gene fragment comprises at least one novel polymorphism identified herein and has a length of at least 10 nucleotides and may range up to the full 30 length of the gene. Preferably, such fragments are between 100 and 3000 nucleotides in length, and more preferably between 200 and 2000 nucleotides in length, and most preferably between 500 and 1000 nucleotides in length.

In describing the EDG6 polymorphic sites identified herein, reference is made to the sense strand of the gene for convenience. However, as recognized by the skilled artisan, nucleic acid 35 molecules containing the EDG6 gene may be complementary double stranded molecules and thus reference to a particular site on the sense strand refers as well to the corresponding site on the complementary antisense strand. Thus, reference may be made to the same polymorphic site on either strand and an oligonucleotide may be designed to hybridize specifically to either strand at a target

region containing the polymorphic site. Thus, the invention also includes single-stranded polynucleotides which are complementary to the sense strand of the EDG6 genomic variants described herein.

5 Polynucleotides comprising a polymorphic gene variant or fragment may be useful for therapeutic purposes. For example, where a patient could benefit from expression, or increased expression, of a particular EDG6 protein isoform, an expression vector encoding the isoform may be administered to the patient. The patient may be one who lacks the EDG6 isogene encoding that isoform or may already have at least one copy of that isogene.

10 In other situations, it may be desirable to decrease or block expression of a particular EDG6 isogene. Expression of an EDG6 isogene may be turned off by transforming a targeted organ, tissue or cell population with an expression vector that expresses high levels of untranslatable mRNA for the isogene. Alternatively, oligonucleotides directed against the regulatory regions (e.g., promoter, introns, enhancers, 3' untranslated region) of the isogene may block transcription. Oligonucleotides targeting the transcription initiation site, e.g., between positions -10 and +10 from the start site are 15 preferred. Similarly, inhibition of transcription can be achieved using oligonucleotides that base-pair with region(s) of the isogene DNA to form triplex DNA (see e.g., Gee et al. in Huber, B.E. and B.I. Carr, Molecular and Immunologic Approaches, Futura Publishing Co., Mt. Kisco, N.Y., 1994). Antisense oligonucleotides may also be designed to block translation of EDG6 mRNA transcribed 20 from a particular isogene. It is also contemplated that ribozymes may be designed that can catalyze the specific cleavage of EDG6 mRNA transcribed from a particular isogene.

25 The oligonucleotides may be delivered to a target cell or tissue by expression from a vector introduced into the cell or tissue *in vivo* or *ex vivo*. Alternatively, the oligonucleotides may be formulated as a pharmaceutical composition for administration to the patient. Oligoribonucleotides and/or oligodeoxynucleotides intended for use as antisense oligonucleotides may be modified to increase stability and half-life. Possible modifications include, but are not limited to phosphorothioate or 2' O-methyl linkages, and the inclusion of nontraditional bases such as inosine and queosine, as well as acetyl-, methyl-, thio-, and similarly modified forms of adenine, cytosine, guanine, thymine, and uracil which are not as easily recognized by endogenous nucleases.

30 The invention also provides an isolated polypeptide comprising a polymorphic variant of the reference EDG6 amino acid sequence shown in Figure 3. The location of a variant amino acid in an EDG6 polypeptide or fragment of the invention is identified by aligning its sequence against SEQ ID NO:3 (Fig. 3). An EDG6 protein variant of the invention comprises an amino acid sequence identical to SEQ ID NO:3 except for having one or more variant amino acids selected from the group consisting of arginine at a position corresponding to amino acid position 155, serine at a position corresponding 35 to amino acid position 164, serine at a position corresponding to amino acid position 189, cysteine at a position corresponding to amino acid position 243, leucine at a position corresponding to amino acid position 365 and methionine at a position corresponding to amino acid position 381. The invention specifically excludes amino acid sequences identical to those previously identified for EDG6,

including SEQ ID NO:3, and previously described fragments thereof. EDG6 protein variants included within the invention comprise all amino acid sequences based on SEQ ID NO:3 and having the combination of amino acid variations described in Table 2 below. In preferred embodiments, an EDG6 protein variant of the invention is encoded by an isogene defined by one of the observed haplotypes 5 shown in Table 5.

Table 2. Novel Polymorphic Variants of EDG6

Polymorphic Variant		Amino Acid Position and Identities					
	Number	155	164	189	243	365	381
5	1	G	G	A	R	R	M
	2	G	G	A	R	L	V
	3	G	G	A	R	L	M
	4	G	G	A	C	R	V
10	5	G	G	A	C	R	M
	6	G	G	A	C	L	V
	7	G	G	A	C	L	M
	8	G	G	S	R	R	V
	9	G	G	S	R	R	M
15	10	G	G	S	R	L	V
	11	G	G	S	R	L	M
	12	G	G	S	C	R	V
	13	G	G	S	C	R	M
	14	G	G	S	C	L	V
20	15	G	G	S	C	L	M
	16	G	S	A	R	R	V
	17	G	S	A	R	R	M
	18	G	S	A	R	L	V
	19	G	S	A	R	L	M
25	20	G	S	A	C	R	V
	21	G	S	A	C	R	M
	22	G	S	A	C	L	V
	23	G	S	A	C	L	M
	24	G	S	S	R	R	V
30	25	G	S	S	R	R	M
	26	G	S	S	R	L	V
	27	G	S	S	R	L	M
	28	G	S	S	C	R	V
	29	G	S	S	C	R	M
35	30	G	S	S	C	L	V
	31	G	S	S	C	L	M
	32	R	G	A	R	R	V

Table 2 Cont. Novel Polymorphic Variants of EDG6

Polymorphic Variant	Number	Amino Acid Position and Identities					
		155	164	189	243	365	381
5	33	R	G	A	R	R	M
	34	R	G	A	R	L	V
	35	R	G	A	R	L	M
	36	R	G	A	C	R	V
10	37	R	G	A	C	R	M
	38	R	G	A	C	L	V
	39	R	G	A	C	L	M
	40	R	G	S	R	R	V
	41	R	G	S	R	R	M
15	42	R	G	S	R	L	V
	43	R	G	S	R	L	M
	44	R	G	S	C	R	V
	45	R	G	S	C	R	M
	46	R	G	S	C	L	V
20	47	R	G	S	C	L	M
	48	R	S	A	R	R	V
	49	R	S	A	R	R	M
	50	R	S	A	R	L	V
	51	R	S	A	R	L	M
25	52	R	S	A	C	R	V
	53	R	S	A	C	R	M
	54	R	S	A	C	L	V
	55	R	S	A	C	L	M
	56	R	S	S	R	R	V
30	57	R	S	S	R	R	M
	58	R	S	S	R	L	V
	59	R	S	S	R	L	M
	60	R	S	S	C	R	V
	61	R	S	S	C	R	M
35	62	R	S	S	C	L	V
	63	R	S	S	C	L	M

The invention also includes EDG6 peptide variants, which are any fragments of an EDG6 protein variant that contain one or more of the amino acid variations shown in Table 2. An EDG6 peptide variant is at least 6 amino acids in length and is preferably any number between 6 and 30 amino acids long, more preferably between 10 and 25, and most preferably between 15 and 20 amino acids long. Such EDG6 peptide variants may be useful as antigens to generate antibodies specific for one of the above EDG6 isoforms. In addition, the EDG6 peptide variants may be useful in drug screening assays.

An EDG6 variant protein or peptide of the invention may be prepared by chemical synthesis or by expressing one of the variant EDG6 genomic and cDNA sequences as described above. Alternatively, the EDG6 protein variant may be isolated from a biological sample of an individual having an EDG6 isogene which encodes the variant protein. Where the sample contains two different EDG6 isoforms (i.e., the individual has different EDG6 isogenes), a particular EDG6 isoform of the invention can be isolated by immunoaffinity chromatography using an antibody which specifically

binds to that particular EDG6 isoform but does not bind to the other EDG6 isoform.

The expressed or isolated EDG6 protein may be detected by methods known in the art, including Coomassie blue staining, silver staining, and Western blot analysis using antibodies specific for the isoform of the EDG6 protein as discussed further below. EDG6 variant proteins can be purified 5 by standard protein purification procedures known in the art, including differential precipitation, molecular sieve chromatography, ion-exchange chromatography, isoelectric focusing, gel electrophoresis, affinity and immunoaffinity chromatography and the like. (Ausubel et. al., 1987, In Current Protocols in Molecular Biology John Wiley and Sons, New York, New York). In the case of immunoaffinity chromatography, antibodies specific for a particular polymorphic variant may be used.

10 A polymorphic variant EDG6 gene of the invention may also be fused in frame with a heterologous sequence to encode a chimeric EDG6 protein. The non-EDG6 portion of the chimeric protein may be recognized by a commercially available antibody. In addition, the chimeric protein may also be engineered to contain a cleavage site located between the EDG6 and non-EDG6 portions so that the EDG6 protein may be cleaved and purified away from the non-EDG6 portion.

15 An additional embodiment of the invention relates to using a novel EDG6 protein isoform in any of a variety of drug screening assays. Such screening assays may be performed to identify agents that bind specifically to all known EDG6 protein isoforms or to only a subset of one or more of these isoforms. The agents may be from chemical compound libraries, peptide libraries and the like. The EDG6 protein or peptide variant may be free in solution or affixed to a solid support. In one 20 embodiment, high throughput screening of compounds for binding to an EDG6 variant may be accomplished using the method described in PCT application WO84/03565, in which large numbers of test compounds are synthesized on a solid substrate, such as plastic pins or some other surface, contacted with the EDG6 protein(s) of interest and then washed. Bound EDG6 protein(s) are then detected using methods well-known in the art.

25 In another embodiment, a novel EDG6 protein isoform may be used in assays to measure the binding affinities of one or more candidate drugs targeting the EDG6 protein.

In yet another embodiment, when a particular EDG6 haplotype or group of EDG6 haplotypes encodes an EDG6 protein variant with an amino acid sequence distinct from that of EDG6 protein 30 isoforms encoded by other EDG6 haplotypes, then detection of that particular EDG6 haplotype or group of EDG6 haplotypes may be accomplished by detecting expression of the encoded EDG6 protein variant using any of the methods described herein or otherwise commonly known to the skilled artisan.

In another embodiment, the invention provides antibodies specific for and immunoreactive with one or more of the novel EDG6 variant proteins described herein. The antibodies may be either 35 monoclonal or polyclonal in origin. The EDG6 protein or peptide variant used to generate the antibodies may be from natural or recombinant sources or produced by chemical synthesis using synthesis techniques known in the art. If the EDG6 protein variant is of insufficient size to be antigenic, it may be conjugated, complexed, or otherwise covalently linked to a carrier molecule to

enhance the antigenicity of the peptide. Examples of carrier molecules, include, but are not limited to, albumins (e.g., human, bovine, fish, ovine), and keyhole limpet hemocyanin (Basic and Clinical Immunology, 1991, Eds. D.P. Stites, and A.I. Terr, Appleton and Lange, Norwalk Connecticut, San Mateo, California).

5 In one embodiment, an antibody specifically immunoreactive with one of the novel protein isoforms described herein is administered to an individual to neutralize activity of the EDG6 isoform expressed by that individual. The antibody may be formulated as a pharmaceutical composition which includes a pharmaceutically acceptable carrier.

10 Antibodies specific for and immunoreactive with one of the novel protein isoforms described herein may be used to immunoprecipitate the EDG6 protein variant from solution as well as react with EDG6 protein isoforms on Western or immunoblots of polyacrylamide gels on membrane supports or substrates. In another preferred embodiment, the antibodies will detect EDG6 protein isoforms in paraffin or frozen tissue sections, or in cells which have been fixed or unfixed and prepared on slides, coverslips, or the like, for use in immunocytochemical, immunohistochemical, and

15 immunofluorescence techniques.

In another embodiment, an antibody specifically immunoreactive with one of the novel EDG6 protein variants described herein is used in immunoassays to detect this variant in biological samples. In this method, an antibody of the present invention is contacted with a biological sample and the formation of a complex between the EDG6 protein variant and the antibody is detected. As described, 20 suitable immunoassays include radioimmunoassay, Western blot assay, immunofluorescent assay, enzyme linked immunoassay (ELISA), chemiluminescent assay, immunohistochemical assay, immunocytochemical assay, and the like (see, e.g., Principles and Practice of Immunoassay, 1991, Eds. Christopher P. Price and David J. Neoman, Stockton Press, New York, New York; Current Protocols in Molecular Biology, 1987, Eds. Ausubel et al., John Wiley and Sons, New York, New York).

25 Standard techniques known in the art for ELISA are described in Methods in Immunodiagnosis, 2nd Ed., Eds. Rose and Bigazzi, John Wiley and Sons, New York 1980; and Campbell et al., 1984, Methods in Immunology, W.A. Benjamin, Inc.). Such assays may be direct, indirect, competitive, or noncompetitive as described in the art (see, e.g., Principles and Practice of Immunoassay, 1991, Eds. Christopher P. Price and David J. Neoman, Stockton Pres, NY, NY; and Oellrich, M., 1984, J. Clin. 30 Chem. Clin. Biochem., 22:895-904). Proteins may be isolated from test specimens and biological samples by conventional methods, as described in Current Protocols in Molecular Biology, supra.

Exemplary antibody molecules for use in the detection and therapy methods of the present invention are intact immunoglobulin molecules, substantially intact immunoglobulin molecules, or those portions of immunoglobulin molecules that contain the antigen binding site. Polyclonal or 35 monoclonal antibodies may be produced by methods conventionally known in the art (e.g., Kohler and Milstein, 1975, Nature, 256:495-497; Campbell Monoclonal Antibody Technology, the Production and Characterization of Rodent and Human Hybridomas, 1985, In: Laboratory Techniques in Biochemistry and Molecular Biology, Eds. Burdon et al., Volume 13, Elsevier Science Publishers, Amsterdam). The

antibodies or antigen binding fragments thereof may also be produced by genetic engineering. The technology for expression of both heavy and light chain genes in *E. coli* is the subject of PCT patent applications, publication number WO 901443, WO 901443 and WO 9014424 and in Huse et al., 1989, *Science*, 246:1275-1281. The antibodies may also be humanized (e.g., Queen, C. et al. 1989 *Proc. 5 Natl. Acad. Sci. USA* 86;10029).

Effect(s) of the polymorphisms identified herein on expression of EDG6 may be investigated by preparing recombinant cells and/or nonhuman recombinant organisms, preferably recombinant animals, containing a polymorphic variant of the EDG6 gene. As used herein, "expression" includes but is not limited to one or more of the following: transcription of the gene into precursor mRNA; 10 splicing and other processing of the precursor mRNA to produce mature mRNA; mRNA stability; translation of the mature mRNA into EDG6 protein (including codon usage and tRNA availability); and glycosylation and/or other modifications of the translation product, if required for proper expression and function.

To prepare a recombinant cell of the invention, the desired EDG6 isogene may be introduced 15 into the cell in a vector such that the isogene remains extrachromosomal. In such a situation, the gene will be expressed by the cell from the extrachromosomal location. In a preferred embodiment, the EDG6 isogene is introduced into a cell in such a way that it recombines with the endogenous EDG6 gene present in the cell. Such recombination requires the occurrence of a double recombination event, thereby resulting in the desired EDG6 gene polymorphism. Vectors for the introduction of genes both 20 for recombination and for extrachromosomal maintenance are known in the art, and any suitable vector or vector construct may be used in the invention. Methods such as electroporation, particle bombardment, calcium phosphate co-precipitation and viral transduction for introducing DNA into cells are known in the art; therefore, the choice of method may lie with the competence and preference of the skilled practitioner. Examples of cells into which the EDG6 isogene may be introduced include, 25 but are not limited to, continuous culture cells, such as COS, NIH/3T3, and primary or culture cells of the relevant tissue type, i.e., they express the EDG6 isogene. Such recombinant cells can be used to compare the biological activities of the different protein variants.

Recombinant nonhuman organisms, i.e., transgenic animals, expressing a variant EDG6 gene 30 are prepared using standard procedures known in the art. Preferably, a construct comprising the variant gene is introduced into a nonhuman animal or an ancestor of the animal at an embryonic stage, i.e., the one-cell stage, or generally not later than about the eight-cell stage. Transgenic animals carrying the constructs of the invention can be made by several methods known to those having skill in the art. One method involves transfecting into the embryo a retrovirus constructed to contain one or 35 more insulator elements, a gene or genes of interest, and other components known to those skilled in the art to provide a complete shuttle vector harboring the insulated gene(s) as a transgene, see e.g., U.S. Patent No. 5,610,053. Another method involves directly injecting a transgene into the embryo. A third method involves the use of embryonic stem cells. Examples of animals into which the EDG6 isogenes may be introduced include, but are not limited to, mice, rats, other rodents, and nonhuman

primates (see "The Introduction of Foreign Genes into Mice" and the cited references therein, In: Recombinant DNA, Eds. J.D. Watson, M. Gilman, J. Witkowski, and M. Zoller; W.H. Freeman and Company, New York, pages 254-272). Transgenic animals stably expressing a human EDG6 isogene and producing human EDG6 protein can be used as biological models for studying diseases related to 5 abnormal EDG6 expression and/or activity, and for screening and assaying various candidate drugs, compounds, and treatment regimens to reduce the symptoms or effects of these diseases.

An additional embodiment of the invention relates to pharmaceutical compositions for treating disorders affected by expression or function of a novel EDG6 isogene described herein. The pharmaceutical composition may comprise any of the following active ingredients: a polynucleotide 10 comprising one of these novel EDG6 isogenes; an antisense oligonucleotide directed against one of the novel EDG6 isogenes, a polynucleotide encoding such an antisense oligonucleotide, or another compound which inhibits expression of a novel EDG6 isogene described herein. Preferably, the composition contains the active ingredient in a therapeutically effective amount. By therapeutically effective amount is meant that one or more of the symptoms relating to disorders affected by 15 expression or function of a novel EDG6 isogene is reduced and/or eliminated. The composition also comprises a pharmaceutically acceptable carrier, examples of which include, but are not limited to, saline, buffered saline, dextrose, and water. Those skilled in the art may employ a formulation most suitable for the active ingredient, whether it is a polynucleotide, oligonucleotide, protein, peptide or 20 small molecule antagonist. The pharmaceutical composition may be administered alone or in combination with at least one other agent, such as a stabilizing compound. Administration of the pharmaceutical composition may be by any number of routes including, but not limited to oral, intravenous, intramuscular, intra-arterial, intramedullary, intrathecal, intraventricular, intradermal, transdermal, subcutaneous, intraperitoneal, intranasal, enteral, topical, sublingual, or rectal. Further 25 details on techniques for formulation and administration may be found in the latest edition of Remington's Pharmaceutical Sciences (Maack Publishing Co., Easton, PA).

For any composition, determination of the therapeutically effective dose of active ingredient and/or the appropriate route of administration is well within the capability of those skilled in the art. For example, the dose can be estimated initially either in cell culture assays or in animal models. The animal model may also be used to determine the appropriate concentration range and route of 30 administration. Such information can then be used to determine useful doses and routes for administration in humans. The exact dosage will be determined by the practitioner, in light of factors relating to the patient requiring treatment, including but not limited to severity of the disease state, general health, age, weight and gender of the patient, diet, time and frequency of administration, other drugs being taken by the patient, and tolerance/response to the treatment.

35 Any or all analytical and mathematical operations involved in practicing the methods of the present invention may be implemented by a computer. In addition, the computer may execute a program that generates views (or screens) displayed on a display device and with which the user can interact to view and analyze large amounts of information relating to the EDG6 gene and its genomic

variation, including chromosome location, gene structure, and gene family, gene expression data, polymorphism data, genetic sequence data, and clinical data population data (e.g., data on ethnogeographic origin, clinical responses, genotypes, and haplotypes for one or more populations). The EDG6 polymorphism data described herein may be stored as part of a relational database (e.g., an instance of an Oracle database or a set of ASCII flat files). These polymorphism data may be stored on the computer's hard drive or may, for example, be stored on a CD-ROM or on one or more other storage devices accessible by the computer. For example, the data may be stored on one or more databases in communication with the computer via a network.

Preferred embodiments of the invention are described in the following examples. Other 10 embodiments within the scope of the claims herein will be apparent to one skilled in the art from consideration of the specification or practice of the invention as disclosed herein. It is intended that the specification, together with the examples, be considered exemplary only, with the scope and spirit of the invention being indicated by the claims which follow the examples.

15 EXAMPLES

The Examples herein are meant to exemplify the various aspects of carrying out the invention and are not intended to limit the scope of the invention in any way. The Examples do not include detailed descriptions for conventional methods employed, such as in the performance of genomic DNA isolation, PCR and sequencing procedures. Such methods are well-known to those skilled in the art 20 and are described in numerous publications, for example, Sambrook, Fritsch, and Maniatis, "Molecular Cloning: A Laboratory Manual", 2nd Edition, Cold Spring Harbor Laboratory Press, USA, (1989).

EXAMPLE 1

This example illustrates examination of various regions of the EDG6 gene for polymorphic 25 sites.

Amplification of Target Regions

The following target regions of the EDG6 gene were amplified using PCR primer pairs. The primers used for each region are represented below by providing the nucleotide positions of their initial 30 and final nucleotides, which correspond to positions in Figure 1.

PCR Primer Pairs

	Fragment No.	Forward Primer	Reverse Primer	PCR Product
35	Fragment 1	3484-3507	complement of 4101-4078	618 nt
	Fragment 2	3698-3717	complement of 4427-4407	730 nt
	Fragment 3	3899-3918	complement of 4509-4491	611 nt
	Fragment 4	3938-3960	complement of 4639-4617	702 nt
	Fragment 5	4266-4286	complement of 4941-4920	676 nt
40	Fragment 6	4625-4646	complement of 5188-5169	564 nt
	Fragment 7	4915-4937	complement of 5583-5563	669 nt
	Fragment 8	5070-5089	complement of 5771-5749	702 nt

These primer pairs were used in PCR reactions containing genomic DNA isolated from immortalized cell lines for each member of the Index Repository. The PCR reactions were carried out 5 under the following conditions:

Reaction volume	= 10 μ l
10 x Advantage 2 Polymerase reaction buffer (Clontech)	= 1 μ l
100 ng of human genomic DNA	= 1 μ l
10 mM dNTP	= 0.4 μ l
10 Advantage 2 Polymerase enzyme mix (Clontech)	= 0.2 μ l
Forward Primer (10 μ M)	= 0.4 μ l
Reverse Primer (10 μ M)	= 0.4 μ l
Water	= 6.6 μ l
15 Amplification profile:	
97°C - 2 min.	1 cycle
97°C - 15 sec.	
70°C - 45 sec.	
20 72°C - 45 sec.	10 cycles
97°C - 15 sec.	
64°C - 45 sec.	
25 72°C - 45 sec.	35 cycles

Sequencing of PCR Products

The PCR products were purified using a Whatman/Polyfiltrronics 100 μ l 384 well unifilter plate essentially according to the manufacturers protocol. The purified DNA was eluted in 50 μ l of 30 distilled water. Sequencing reactions were set up using Applied Biosystems Big Dye Terminator chemistry essentially according to the manufacturers protocol. The purified PCR products were sequenced in both directions using the primer sets described previously or those represented below by the nucleotide positions of their initial and final nucleotides, which correspond to positions in Figure 1. Reaction products were purified by isopropanol precipitation, and run on an Applied Biosystems 3700 35 DNA Analyzer.

Sequencing Primer Pairs

Fragment No.	Forward Primer	Reverse Primer
Fragment 1	3516-3534	complement of 4059-4041
Fragment 2	3757-3776	complement of 4321-4302
Fragment 3	3914-3932	complement of 4386-4368
Fragment 4	4071-4090	complement of 4608-4589
Fragment 5	4403-4423	complement of 4875-4857
Fragment 6	4670-4689	complement of 5152-5134
45 Fragment 7	4939-4958	complement of 5428-5410
Fragment 8	5149-5168	complement of 5691-5671

Analysis of Sequences for Polymorphic Sites

Sequence information for a minimum of 80 humans was analyzed for the presence of polymorphisms using the Polyphred program (Nickerson et al., *Nucleic Acids Res.* 14:2745-2751, 1997). The presence of a polymorphism was confirmed on both strands. The polymorphisms and their 5 locations in the EDG6 gene are listed in Table 3 below.

Table 3. Polymorphic Sites Identified in the EDG6 Gene

	Polymorphic Site Number	PolyId ^a	Nucleotide Position	Reference Allele	Variant Allele	CDS Position	Variant AA
10	PS1	3216843	3591	G	A		
	PS2	3216845	3697	C	T		
	PS3	3216847	3804	C	T		
	PS4	3216851	3818	A	G		
15	PS5	3216859	4123	C	T	114	R38R
	PS6	3216861	4240	G	A	231	S77S
	PS7	3216863	4472	G	A	463	G155R
	PS8	3216865	4499	G	A	490	G164S
	PS9	3216867	4531	G	A	522	A174A
20	PS10	3216869	4574	G	T	565	A189S
	PS11	3216871	4736	C	T	727	R243C
	PS12	3216873	4813	C	T	804	F268F
	PS13	3216877	5068	C	T	1059	S353S
	PS14	3216879	5103	G	T	1094	R365L
25	PS15	3216883	5150	G	A	1141	V381M
	PS16	3216885	5179	G	A		
	PS17	3216887	5301	G	A		
	PS18	3216889	5333	G	A		
	PS19	3216893	5448	G	C		
30	PS20	3216895	5560	G	A		
	PS21	3216899	5580	G	A		
	PS22	3216901	5587	C	T		
	PS23	3216903	5606	G	C		

^aPolyId is a unique identifier assigned to each PS by Genaissance Pharmaceuticals, Inc.

35

EXAMPLE 2

This example illustrates analysis of the EDG6 polymorphisms identified in the Index Repository for human genotypes and haplotypes.

The different genotypes containing these polymorphisms that were observed in the reference 40 population are shown in Table 4 below, with the haplotype pair indicating the combination of haplotypes determined for the individual using the haplotype derivation protocol described below. In Table 4, homozygous positions are indicated by one nucleotide and heterozygous positions are indicated by two nucleotides. Missing nucleotides in any given genotype in Table 4 were inferred based on linkage disequilibrium and/or Mendelian inheritance.

Table 4 (Part1). Genotypes and Haplotype Pairs Observed for EDG6 Gene

Genotype Number	Polymorphic Sites										HAP	Pair
	PS1	PS2	PS3	PS4	PS5	PS6	PS7	PS8	PS9	PS10		
5	1	G	C	C	G	C	G	G	G	G	18	18
	2	G	C	C	G	C	G	G	G	G	17	17
	3	G	C	C	A	C	G	G	G	G	5	5
	4	G	C	C	G	C	G	G	G	G	16	16
10	5	G	C/T	C	G/A	C	G	G	G	G	17	24
	6	G	C	C	A	C	G	G	G	G	5	7
	7	G	C	C	G	C	G	G/A	G	G	17	9
	8	G	C	C	G	C	G	G	G	G	17	20
	9	G	C	C	G	C/T	G	G	G	G	17	22
15	10	G/A	C	C	G	C	G	G	G	G	17	1
	11	G	C	C	G	C	G	G	G/A	G	17	10
	12	G	C	C	G/A	C	G	G	G	G	17	6
	13	G	C	C	G	C	G	G	G	G	17	12
	14	G	C	C	G/A	C	G	G	G	G	17	7
20	15	G	C	C	G	C	G	G	G	G	17	13
	16	G	C	C	G	C	G	G	G/A	G	18	10
	17	G	C	C	A	C	G/A	G	G	G	5	3
	18	G	C	C	G/A	C	G/A	G	G	G	17	3
	19	G	C	C	A	C	G	G	G	G	5	6
25	20	G	C	C	G	C	G/A	G	G	G	17	8
	21	G	C	C	G	C	G	G	G	G	18	14
	22	G	C	C	G	C	G	G	G	G	17	14
	23	G	C	C	G/A	C	G	G	G	G	17	5
	24	G	C	C	G	C	G	G	G	G	17	15
30	25	G	C	C	C/T	G	C	G	G	G	17	23
	26	G	C	C	G	C	G	G	G	G/A	17	11
	27	G	C	C	A	C	G	G	G	G	5	4
	28	G	C	C	G/A	C	G	G	G	G	18	6
	29	G	C	C	G	C	G	G	G	G	17	18
35	30	G	C	C	G	C	G	G	G	G	17	16
	31	G	C	C	G/A	C	G/A	G	G	G	18	3
	32	G	C	C	G	C	G	G	G	G/T	17	21
	33	G	C	C	G/A	C	G	G	G	G	18	5
	34	G/A	C	C	G	C	G	G	G	G	17	2
40	35	G	C	C	G	C	G	G	G	G	17	19

Table 4 (Part2). Genotypes and Haplotype Pairs Observed for EDG6 Gene

Genotype	Polymorphic Sites										HAP	Pair
Number	PS11	PS12	PS13	PS14	PS15	PS16	PS17	PS18	PS19	PS20		
5	1	C	C	C	G	G	G	G	G	G	18	18
	2	C	C	C	G	G	G	G	G	G	17	17
	3	C	C	C	G	G	G	G	G	G	5	5
	4	C	C	C	G	G	G	G	G	G	16	16
	5	C	C	C	G/T	G	G	G	G	G	17	24
	6	C/T	C	C	G	G	G	G	G	G	5	7
10	7	C	C	C	G	G	G	G	G	G	17	9
	8	C	C	C/T	G	G	G	G	G	G	17	20
	9	C	C	C	G	G	G	G	G	G	17	22
	10	C	C	C	G	G	G	G	G	G	17	1
	11	C	C	C	G	G	G	G	G	G	17	10
	12	C	C	C	G	G	G	G	G	G	17	6
15	13	C	C	C	G	G/A	G	G	G	G	17	12
	14	C/T	C	C	G	G	G	G	G	G	17	7
	15	C	C	C	G	G	G	G	G/A	G	17	13
	16	C	C	C	G	G	G	G	G	G	18	10
	17	C	C	C	G	G	G	G	G	G	5	3
	18	C	C	C	G	G	G	G	G	G	17	3
20	19	C	C	C	G	G	G	G	G	G	5	6
	20	C	C	C	G	G	G	G	G	G	17	8
	21	C	C	C	G	G	G	G	G	G/C	18	14
	22	C	C	C	G	G	G	G	G	G/C	17	14
	23	C	C	C	G	G	G	G	G	G	17	5
	24	C	C	C	G	G	G	G	G	G/A	17	15
25	25	C	C	C	G	G	G	G	G	G	17	23
	26	C/T	C	C	G	G	G	G	G	G	17	11
	27	C	C	C	G	G	G	G/A	G	G	5	4
	28	C	C	C	G	G	G	G	G	G	18	6
	29	C	C	C	G	G	G	G	G	G	17	18
	30	C	C	C	G	G	G	G	G	G	17	16
35	31	C	C	C	G	G	G	G	G	G	18	3
	32	C	C	C	G	G	G	G	G	G	17	21
	33	C	C	C	G	G	G	G	G	G	18	5
	34	C	C	C	G	G	G	G	G	G	17	2
	35	C	C	C/T	G	G	G/A	G	G	G	17	19

40 Table 4 (Part3). Genotypes and Haplotype Pairs Observed for EDG6 Gene

Genotype	Polymorphic Sites			HAP	Pair	
Number	PS21	PS22	PS23	HAP	Pair	
45	1	G	T	G	18	18
	2	G	C	G	17	17
	3	G	C	G	5	5
	4	A	C	G	16	16
	5	G	C	G	17	24
	6	G	C/T	G	5	7
50	7	G	C/T	G	17	9
	8	G	C	G	17	20
	9	G	C	G	17	22
	10	G	C	G	17	1
	11	G	C	G	17	10
	12	G	C/T	G	17	6
55	13	G	C	G	17	12
	14	G	C/T	G	17	7
	15	G	C	G	17	13

Table 4 (Part3 cont.). Genotypes and Haplotype Pairs Observed for EDG6 Gene

Genotype Polymorphic Sites
Number PS21 PS22 PS23 HAP Pair

5	16	G	T/C	G	18	10
	17	G	C/T	G	5	3
	18	G	C/T	G	17	3
	19	G	C/T	G	5	6
	20	G	C	G	17	8
10	21	G	T/C	G	18	14
	22	G	C	G	17	14
	23	G	C	G	17	5
	24	G	C	G	17	15
	25	G	C/T	G	17	23
15	26	G	C/T	G	17	11
	27	G	C	G	5	4
	28	G	T	G	18	6
	29	G	C/T	G	17	18
	30	G/A	C	G	17	16
20	31	G	T	G	18	3
	32	G	C	G	17	21
	33	G	T/C	G	18	5
	34	G	C/T	G/C	17	2
	35	G	C	G	17	19

25

The haplotype pairs shown in Table 4 were estimated from the unphased genotypes using a computer-implemented extension of Clark's algorithm (Clark, A.G. 1990 *Mol Bio Evol* 7, 111-122) for assigning haplotypes to unrelated individuals in a population sample, as described in U.S. Provisional Application Serial No. 60/198,340 entitled "A Method and System for Determining Haplotypes from a Collection of Polymorphisms" and the corresponding International Application, PCT/US01/12831. In this method, haplotypes are assigned directly from individuals who are homozygous at all sites or heterozygous at no more than one of the variable sites. This list of haplotypes is then used to deconvolute the unphased genotypes in the remaining (multiply heterozygous) individuals. In our analysis, the list of haplotypes was augmented with haplotypes obtained from two families (one three-generation Caucasian family and one two-generation African-American family).

30 By following this protocol, it was determined that the Index Repository examined herein and, by extension, the general population contains the 24 human EDG6 haplotypes shown in Table 5 below.

An EDG6 isogene defined by a full-haplotype shown in Table 5 below comprises the regions of the SEQ ID NOS indicated in Table 5, with their corresponding set of polymorphic locations and 40 identities, which are also set forth in Table 5.

Table 5 (Part A). Haplotypes of the EDG6 Gene

	Haplotype Number ^a										PS ^b	PS	Seq ID	Region
	1	2	3	4	5	6	7	8	9	10	No.	Position ^c	No. ^d	Examined ^e
5	A	A	G	G	G	G	G	G	G	G	1	3591	1/119	3484-5771
	C	C	C	C	C	C	C	C	C	C	2	3697	1/119	3484-5771
	C	C	C	C	C	C	C	C	C	C	3	3804	1/119	3484-5771
	G	G	A	A	A	A	A	G	G	G	4	3818	1/119	3484-5771
	C	C	C	C	C	C	C	C	C	C	5	4123	1/119	3484-5771
10	G	G	A	G	G	G	G	A	G	G	6	4240	1/119	3484-5771
	G	G	G	G	G	G	G	G	A	G	7	4472	1/119	3484-5771
	G	G	G	G	G	G	G	G	G	A	8	4499	1/119	3484-5771
	G	G	G	G	G	G	G	G	G	G	9	4531	1/119	3484-5771
	G	G	G	G	G	G	G	G	G	G	10	4574	1/119	3484-5771
15	C	C	C	C	C	C	T	C	C	C	11	4736	1/119	3484-5771
	C	C	C	C	C	C	C	C	C	C	12	4813	1/119	3484-5771
	C	C	C	C	C	C	C	C	C	C	13	5068	1/119	3484-5771
	G	G	G	G	G	G	G	G	G	G	14	5103	1/119	3484-5771
	G	G	G	G	G	G	G	G	G	G	15	5150	1/119	3484-5771
20	G	G	G	G	G	G	G	G	G	G	16	5179	1/119	3484-5771
	G	G	G	A	G	G	G	G	G	G	17	5301	1/119	3484-5771
	G	G	G	G	G	G	G	G	G	G	18	5333	1/119	3484-5771
	G	G	G	G	G	G	G	G	G	G	19	5448	1/119	3484-5771
	G	G	G	G	G	G	G	G	G	G	20	5560	1/119	3484-5771
25	G	G	G	G	G	G	G	G	G	G	21	5580	1/119	3484-5771
	C	T	T	C	C	T	T	C	T	C	22	5587	1/119	3484-5771
	G	C	G	G	G	G	G	G	G	G	23	5606	1/119	3484-5771

Table 5 (Part B). Haplotypes of the EDG6 Gene

	Haplotype Number ^a										PS ^b	PS	Seq ID	Region
	11	12	13	14	15	16	17	18	19	20	No.	Pos. ^c	No. ^d	Examined ^e
30	G	G	G	G	G	G	G	G	G	G	1	3591	1/119	3484-5771
	C	C	C	C	C	C	C	C	C	C	2	3697	1/119	3484-5771
	C	C	C	C	C	C	C	C	C	C	3	3804	1/119	3484-5771
	G	G	G	G	G	G	G	G	G	G	4	3818	1/119	3484-5771
	C	C	C	C	C	C	C	C	C	C	5	4123	1/119	3484-5771
35	G	G	G	G	G	G	G	G	G	G	6	4240	1/119	3484-5771
	G	G	G	G	G	G	G	G	G	G	7	4472	1/119	3484-5771
	C	C	C	C	C	C	C	C	C	C	8	4499	1/119	3484-5771
	A	G	G	G	G	G	G	G	G	G	9	4531	1/119	3484-5771
	G	G	G	G	G	G	G	G	G	G	10	4574	1/119	3484-5771
40	G	G	G	G	G	G	G	G	G	G	11	4736	1/119	3484-5771
	C	C	C	C	C	C	C	C	C	C	12	4813	1/119	3484-5771
	T	C	C	C	C	C	C	C	C	C	13	5068	1/119	3484-5771
	G	G	G	G	G	G	G	G	G	G	14	5103	1/119	3484-5771
	G	A	G	G	G	G	G	G	G	G	15	5150	1/119	3484-5771
45	G	G	G	G	G	G	G	G	G	G	16	5179	1/119	3484-5771
	G	G	G	G	G	G	G	G	G	G	17	5301	1/119	3484-5771
	G	A	G	G	G	G	G	G	G	G	18	5333	1/119	3484-5771
	G	G	G	G	G	G	G	G	G	G	19	5448	1/119	3484-5771
	G	G	G	G	G	G	G	G	G	G	20	5560	1/119	3484-5771
50	G	G	A	G	G	G	G	G	G	G	21	5580	1/119	3484-5771
	G	G	G	C	G	G	G	G	G	G	22	5587	1/119	3484-5771
	T	C	C	C	C	C	C	T	C	C	23	5606	1/119	3484-5771
	G	G	G	G	G	G	G	G	G	G	24	5606	1/119	3484-5771
	G	G	G	G	G	G	G	G	G	G	25	5606	1/119	3484-5771
55	G	G	G	G	G	G	G	G	G	G	26	5606	1/119	3484-5771

Table 5 (Part C). Haplotypes of the EDG6 Gene

	Haplotype Number ^a				PS ^b	PS No.	Seq ID Position ^c	No. ^d	Region
5	21	22	23	24	G	1	3591	1/119	3484-5771
	C	C	C	T	C	2	3697	1/119	3484-5771
	C	C	T	C	G	3	3804	1/119	3484-5771
	G	G	G	A	G	4	3818	1/119	3484-5771
	C	T	C	C	T	5	4123	1/119	3484-5771
	G	G	G	G	G	6	4240	1/119	3484-5771
10	G	G	G	G	G	7	4472	1/119	3484-5771
	G	G	G	G	G	8	4499	1/119	3484-5771
	G	G	G	G	G	9	4531	1/119	3484-5771
	T	G	G	G	G	10	4574	1/119	3484-5771
	C	C	C	C	C	11	4736	1/119	3484-5771
	C	C	C	C	C	12	4813	1/119	3484-5771
15	C	C	C	C	C	13	5068	1/119	3484-5771
	G	G	G	T	T	14	5103	1/119	3484-5771
	G	G	G	G	G	15	5150	1/119	3484-5771
	G	G	G	G	G	16	5179	1/119	3484-5771
	G	G	G	G	G	17	5301	1/119	3484-5771
	G	G	G	G	G	18	5333	1/119	3484-5771
20	G	G	G	G	G	19	5448	1/119	3484-5771
	G	G	G	G	G	20	5560	1/119	3484-5771
	G	G	G	G	G	21	5580	1/119	3484-5771
	C	C	T	C	T	22	5587	1/119	3484-5771
	G	G	G	G	G	23	5606	1/119	3484-5771

^aAlleles for EDG6 haplotypes are presented 5' to 3' in each column

^bPS = polymorphic site;

^cPosition of PS within the indicated SEQ ID NO, with the 1st position number referring to the first SEQ ID NO and the 2nd position number referring to the 2nd SEQ ID NO;

^d1st SEQ ID NO refers to Figure 1, with the two alternative allelic variants of each polymorphic site indicated by the appropriate nucleotide symbol; 2nd SEQ ID NO is a modified version of the 1st SEQ ID NO that comprises the context sequence of each polymorphic site, PS1-PS23, to facilitate electronic searching of the haplotypes;

^eRegion examined represents the nucleotide positions defining the start and stop positions within the 1st SEQ ID NO of the sequenced region.

SEQ ID NO:1 refers to Figure 1, with the two alternative allelic variants of each polymorphic site indicated by the appropriate nucleotide symbol. SEQ ID NO:119 is a modified version of SEQ ID NO:1 that shows the context sequence of each of PS1-PS23 in a uniform format to facilitate electronic searching of the EDG6 haplotypes. For each polymorphic site, SEQ ID NO:119 contains a block of 60 bases of the nucleotide sequence encompassing the centrally-located polymorphic site at the 30th position, followed by 60 bases of unspecified sequence to represent that each polymorphic site is separated by genomic sequence whose composition is defined elsewhere herein.

Table 6 below shows the percent of chromosomes characterized by a given EDG6 haplotype for all unrelated individuals in the Index Repository for which haplotype data was obtained. The percent of these unrelated individuals who have a given EDG6 haplotype pair is shown in Table 7. In Tables 6 and 7, the "Total" column shows this frequency data for all of these unrelated individuals, while the other columns show the frequency data for these unrelated individuals categorized according

to their self-identified ethnogeographic origin. Abbreviations used in Tables 6 and 7 are AF = African Descent, AS = Asian, CA = Caucasian, HL = Hispanic-Latino, and AM = Native American.

Table 6. Frequency of Observed EDG6 Haplotypes In Unrelated Individuals

5

	HAP No.	HAP ID	Total	CA	AF	AS	HL	AM
	1	3219666	0.61	2.38	0.0	0.0	0.0	0.0
	2	3219662	0.61	0.0	0.0	2.5	0.0	0.0
	3	3219649	2.44	4.76	0.0	0.0	5.56	0.0
10	4	3219663	0.61	2.38	0.0	0.0	0.0	0.0
	5	3219646	12.8	19.05	7.5	0.0	22.22	33.33
	6	3219647	3.05	9.52	0.0	0.0	2.78	0.0
	7	3219652	1.22	2.38	0.0	0.0	2.78	0.0
	8	3219653	1.22	4.76	0.0	0.0	0.0	0.0
15	9	3219665	0.61	0.0	0.0	0.0	0.0	16.67
	10	3219654	1.22	0.0	0.0	5.0	0.0	0.0
	11	3219661	0.61	0.0	0.0	2.5	0.0	0.0
	12	3219669	0.61	0.0	0.0	0.0	2.78	0.0
	13	3219673	0.61	0.0	2.5	0.0	0.0	0.0
20	14	3219650	2.44	0.0	10.0	0.0	0.0	0.0
	15	3219677	0.61	0.0	0.0	2.5	0.0	0.0
	16	3219648	3.66	0.0	0.0	15.0	0.0	0.0
	17	3219644	41.46	30.95	50.0	47.5	36.11	50.0
	18	3219645	21.95	23.81	22.5	20.0	25.0	0.0
25	19	3219655	0.61	0.0	2.5	0.0	0.0	0.0
	20	3219678	0.61	0.0	2.5	0.0	0.0	0.0
	21	3219676	0.61	0.0	0.0	2.5	0.0	0.0
	22	3219675	0.61	0.0	0.0	2.5	0.0	0.0
	23	3219664	0.61	0.0	2.5	0.0	0.0	0.0
30	24	3219660	0.61	0.0	0.0	0.0	2.78	0.0

Table 7. Frequency of Observed EDG6 Haplotype Pairs In Unrelated Individuals

HAP1	HAP2	Total	CA	AF	AS	HL	AM
18	18	8.54	14.29	5.0	10.0	5.56	0.0
5	17	13.41	9.52	20.0	15.0	5.56	33.33
5	5	3.66	4.76	0.0	0.0	5.56	33.33
16	16	1.22	0.0	0.0	5.0	0.0	0.0
17	24	1.22	0.0	0.0	0.0	5.56	0.0
5	7	1.22	0.0	0.0	0.0	5.56	0.0
10	17	9	1.22	0.0	0.0	0.0	33.33
	17	20	1.22	0.0	5.0	0.0	0.0
	17	22	1.22	0.0	0.0	5.0	0.0
	17	1	1.22	4.76	0.0	0.0	0.0
	17	10	1.22	0.0	0.0	5.0	0.0
15	17	6	1.22	4.76	0.0	0.0	0.0
	17	12	1.22	0.0	0.0	0.0	5.56
	17	7	1.22	4.76	0.0	0.0	0.0
	17	13	1.22	0.0	5.0	0.0	0.0
	18	10	1.22	0.0	0.0	5.0	0.0
20	5	3	1.22	0.0	0.0	0.0	5.56
	17	3	2.44	4.76	0.0	0.0	5.56
	5	6	3.66	9.52	0.0	0.0	5.56
	17	8	2.44	9.52	0.0	0.0	0.0
	18	14	2.44	0.0	10.0	0.0	0.0
25	17	14	2.44	0.0	10.0	0.0	0.0
	17	5	7.32	9.52	10.0	0.0	11.11
	17	15	1.22	0.0	0.0	5.0	0.0
	17	23	1.22	0.0	5.0	0.0	0.0
	17	11	1.22	0.0	0.0	5.0	0.0
30	5	4	1.22	4.76	0.0	0.0	0.0
	18	6	1.22	4.76	0.0	0.0	0.0
	17	18	17.07	4.76	20.0	15.0	33.33
	17	16	4.88	0.0	0.0	20.0	0.0
	18	3	1.22	4.76	0.0	0.0	0.0
35	17	21	1.22	0.0	0.0	5.0	0.0
	18	5	3.66	4.76	5.0	0.0	5.56
	17	2	1.22	0.0	0.0	5.0	0.0
	17	19	1.22	0.0	5.0	0.0	0.0

40

The size and composition of the Index Repository were chosen to represent the genetic diversity across and within four major population groups comprising the general United States population. For example, as described in Table 1 above, this repository contains approximately equal sample sizes of African-descent, Asian-American, European-American, and Hispanic-Latino population groups. Almost all individuals representing each group had all four grandparents with the same ethnogeographic background. The number of unrelated individuals in the Index Repository provides a sample size that is sufficient to detect SNPs and haplotypes that occur in the general population with high statistical certainty. For instance, a haplotype that occurs with a frequency of 5% in the general population has a probability higher than 99.9% of being observed in a sample of 80 individuals from the general population. Similarly, a haplotype that occurs with a frequency of 10% in a specific population group has a 99% probability of being observed in a sample of 20 individuals from

that population group. In addition, the size and composition of the Index Repository means that the relative frequencies determined therein for the haplotypes and haplotype pairs of the EDG6 gene are likely to be similar to the relative frequencies of these EDG6 haplotypes and haplotype pairs in the general U.S. population and in the four population groups represented in the Index Repository. The 5 genetic diversity observed for the three Native Americans is presented because it is of scientific interest, but due to the small sample size it lacks statistical significance.

In view of the above, it will be seen that the several advantages of the invention are achieved and other advantageous results attained.

As various changes could be made in the above methods and compositions without departing 10 from the scope of the invention, it is intended that all matter contained in the above description and shown in the accompanying drawings shall be interpreted as illustrative and not in a limiting sense.

All references cited in this specification, including patents and patent applications, are hereby incorporated in their entirety by reference. The discussion of references herein is intended merely to summarize the assertions made by their authors and no admission is made that any reference 15 constitutes prior art. Applicants reserve the right to challenge the accuracy and pertinency of the cited references.

What is Claimed is:

1. A method for haplotyping the endothelial differentiation, G-protein-coupled receptor 6 (EDG6) gene of an individual, which comprises determining which of the EDG6 haplotypes shown in the table immediately below defines one copy of the individual's EDG6 gene,
 5 wherein each of the EDG6 haplotypes comprises a set of polymorphisms whose locations and identities are set forth in the table immediately below:

Haplotype Number ^a												PS ^b	PS
	1	2	3	4	5	6	7	8	9	10	Number	Position ^c	
10	A	A	G	G	G	G	G	G	G	G	1	3591	
	C	C	C	C	C	C	C	C	C	C	2	3697	
	C	C	C	C	C	C	C	C	C	C	3	3804	
	G	G	A	A	A	A	A	G	G	G	4	3818	
	C	C	C	C	C	C	C	C	C	C	5	4123	
	G	G	A	G	G	G	G	A	G	G	6	4240	
15	G	G	G	G	G	G	G	A	G	G	7	4472	
	G	G	G	G	G	G	G	G	A	G	8	4499	
	G	G	G	G	G	G	G	G	G	A	9	4531	
	G	G	G	G	G	G	G	G	G	G	10	4574	
	C	C	C	C	C	C	T	C	C	C	11	4736	
	C	C	C	C	C	C	C	C	C	C	12	4813	
20	C	C	C	C	C	C	C	C	C	C	13	5068	
	G	G	G	G	G	G	G	G	G	G	14	5103	
	G	G	G	G	G	G	G	G	G	G	15	5150	
	G	G	G	G	G	G	G	G	G	G	16	5179	
	G	G	G	A	G	G	G	G	G	G	17	5301	
	G	G	G	G	G	G	G	G	G	G	18	5333	
25	G	G	G	G	G	G	G	G	G	G	19	5448	
	G	G	G	G	G	G	G	G	G	G	20	5560	
	G	G	G	G	G	G	G	G	G	G	21	5580	
	C	T	T	C	C	T	T	C	T	C	22	5587	
	G	C	G	G	G	G	G	G	G	G	23	5606	

Haplotype Number ^a												PS ^b	PS
	11	12	13	14	15	16	17	18	19	20	Number	Position ^c	
5	G	G	G	G	G	G	G	G	G	G	1	3591	
	C	C	C	C	C	C	C	C	C	C	2	3697	
	C	C	C	C	C	C	C	C	C	C	3	3804	
	G	G	G	G	G	G	G	G	G	G	4	3818	
	C	C	C	C	C	C	C	C	C	C	5	4123	
	G	G	G	G	G	G	G	G	G	G	6	4240	
	G	G	G	G	G	G	G	G	G	G	7	4472	
10	G	G	G	G	G	G	G	G	G	G	8	4499	
	A	G	G	G	G	G	G	G	G	G	9	4531	
	G	G	G	G	G	G	G	G	G	G	10	4574	
	C	C	C	C	C	C	C	C	C	C	11	4736	
	T	C	C	C	C	C	C	C	C	C	12	4813	
15	C	C	C	C	C	C	C	C	T	T	13	5068	
	G	G	G	G	G	G	G	G	G	G	14	5103	
	G	A	G	G	G	G	G	G	G	G	15	5150	
	G	G	G	G	G	G	G	G	A	G	16	5179	
	G	G	G	G	G	G	G	G	G	G	17	5301	
20	G	G	A	G	G	G	G	G	G	G	18	5333	
	G	G	G	C	G	G	G	G	G	G	19	5448	
	G	G	G	G	A	G	G	G	G	G	20	5560	
	G	G	G	G	G	A	G	G	G	G	21	5580	
	T	C	C	C	C	C	C	T	C	C	22	5587	
25	G	G	G	G	G	G	G	G	G	G	23	5606	

Haplotype Number ^a				PS ^b	PS
	21	22	23	Number	Position ^c
30	G	G	G	1	3591
	C	C	C	2	3697
	C	C	T	3	3804
	G	G	G	4	3818
	C	T	C	5	4123
	G	G	G	6	4240
	G	G	G	7	4472
35	G	G	G	8	4499
	G	G	G	9	4531
	T	G	G	10	4574
	C	C	C	11	4736
	C	C	C	12	4813
40	C	C	C	13	5068
	G	G	G	14	5103
	G	G	G	15	5150
	G	G	G	16	5179
	G	G	G	17	5301
45	G	G	G	18	5333
	G	G	G	19	5448
	G	G	G	20	5560
	G	G	G	21	5580
	C	C	T	22	5587
50	G	G	G	23	5606

^aAlleles for haplotypes are presented 5' to 3' in each column

^bPS = polymorphic site;

^cPosition of PS within SEQ ID NO:1.

2. The method of claim 1, wherein the determining step comprises identifying the phased

sequence of nucleotides present at each of PS1-PS23 on the one copy of the individual's EDG6 gene.

3. A method for haplotyping the endothelial differentiation, G-protein-coupled receptor 6 (EDG6) gene of an individual, which comprises determining which of the EDG6 haplotype pairs shown in the table immediately below defines both copies of the individual's EDG6 gene, wherein each of the EDG6 haplotype pairs consists of first and second haplotypes which comprise first and second sets of polymorphisms whose locations and identities are set forth in the table immediately below:

Haplotype Pair ^a										PS ^b	PS
	18/18	17/17	5/5	16/16	17/24	5/7	17/9	17/20	Number	Position ^c	
10	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	1	3591	
	C/C	C/C	C/C	C/C	C/T	C/C	C/C	C/C	2	3697	
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	3	3804	
15	G/G	G/G	A/A	G/G	G/A	A/A	G/G	G/G	4	3818	
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	5	4123	
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	6	4240	
	G/G	G/G	G/G	G/G	G/G	G/G	G/A	G/G	7	4472	
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	8	4499	
20	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	9	4531	
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	10	4574	
	C/C	C/C	C/C	C/C	C/T	C/C	C/C	C/C	11	4736	
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	12	4813	
25	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/T	13	5068	
	G/G	G/G	G/G	G/G	G/T	G/G	G/G	G/G	14	5103	
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	15	5150	
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	16	5179	
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	17	5301	
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	18	5333	
30	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	19	5448	
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	20	5560	
	G/G	G/G	G/G	A/A	G/G	G/G	G/G	G/G	21	5580	
	T/T	C/C	C/C	C/C	C/T	C/T	C/C	C/C	22	5587	
35	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	23	5606	

	Haplotype Pair ^a							PS ^b	PS	
	17/22	17/1	17/10	17/6	17/12	17/7	17/13	18/10	Number	Position ^c
5	G/G	G/A	G/G	G/G	G/G	G/G	G/G	G/G	1	3591
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	2	3697
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	3	3804
	G/G	G/G	G/G	G/A	G/G	G/A	G/G	G/G	4	3818
	C/T	C/C	C/C	C/C	C/C	C/C	C/C	C/C	5	4123
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	6	4240
10	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	7	4472
	G/G	G/G	G/A	G/G	G/G	G/G	G/G	G/A	8	4499
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	9	4531
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	10	4574
	C/C	C/C	C/C	C/C	C/T	C/T	C/C	C/C	11	4736
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	12	4813
15	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	13	5068
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	14	5103
	G/G	G/G	G/G	G/A	G/G	G/G	G/G	G/G	15	5150
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	16	5179
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	17	5301
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/A	18	5333
20	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	19	5448
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	20	5560
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	21	5580
	C/C	C/C	C/C	C/T	C/C	C/T	C/C	T/C	22	5587
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	23	5606

	Haplotype Pair ^a							PS ^b	PS	
	5/3	17/3	5/6	17/8	18/14	17/14	17/5	17/15	Number	Position ^c
30	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	1	3591
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	2	3697
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	3	3804
	A/A	A/A	A/A	G/G	G/G	G/G	G/A	G/G	4	3818
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	5	4123
	G/A	G/A	G/A	G/A	G/G	G/G	G/G	G/G	6	4240
35	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	7	4472
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	8	4499
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	9	4531
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	10	4574
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	11	4736
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	12	4813
40	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	13	5068
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	14	5103
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	15	5150
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	16	5179
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	17	5301
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	18	5333
45	G/G	G/G	G/G	G/G	G/C	G/C	G/G	G/G	19	5448
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/A	20	5560
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	21	5580
	C/T	C/T	C/T	C/C	T/C	C/C	C/C	C/C	22	5587
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	23	5606

	Haplotype	Pair ^a	PS ^b	PS			
	17/23	17/11 5/4	18/6 17/18	17/16 18/3	17/21	Number	Position ^c
5	G/G	G/G G/G	G/G G/G	G/G G/G	G/G G/G	1	3591
	C/C	C/C C/C	C/C C/C	C/C C/C	C/C C/C	2	3697
	C/T	C/C C/C	C/C C/C	C/C C/C	C/C C/C	3	3804
	G/G	G/G A/A	G/A G/G	G/G G/A	G/A G/G	4	3818
	C/C	C/C C/C	C/C C/C	C/C C/C	C/C C/C	5	4123
	G/G	G/G G/G	G/G G/G	G/G G/A	G/G G/G	6	4240
	G/G	G/G G/G	G/G G/G	G/G G/G	G/G G/G	7	4472
10	G/G	G/G G/G	G/G G/G	G/G G/G	G/G G/G	8	4499
	G/G	G/G G/A	G/G G/G	G/G G/G	G/G G/G	9	4531
	G/G	G/G G/G	G/G G/G	G/G G/G	G/T G/G	10	4574
	C/C	C/C C/C	C/C C/C	C/C C/C	C/C C/C	11	4736
	C/C	C/T C/C	C/C C/C	C/C C/C	C/C C/C	12	4813
15	C/C	C/C C/C	C/C C/C	C/C C/C	C/C C/C	13	5068
	G/G	G/G G/G	G/G G/G	G/G G/G	G/G G/G	14	5103
	G/G	G/G G/G	G/G G/G	G/G G/G	G/G G/G	15	5150
	G/G	G/G G/G	G/G G/G	G/G G/G	G/G G/G	16	5179
	G/G	G/G G/A	G/G G/G	G/G G/G	G/G G/G	17	5301
20	G/G	G/G G/G	G/G G/G	G/G G/G	G/G G/G	18	5333
	G/G	G/G G/G	G/G G/G	G/G G/G	G/G G/G	19	5448
	G/G	G/G G/G	G/G G/G	G/G G/G	G/G G/G	20	5560
	G/G	G/G G/G	G/G G/G	G/A G/G	G/G G/G	21	5580
	C/T	C/T C/C	T/T C/T	C/C T/T	C/C C/T	22	5587
25	G/G	G/G G/G	G/G G/G	G/G G/G	G/G G/G	23	5606

	Haplotype	Pair ^a	PS ^b	PS
	18/5	17/2 17/19	Number	Position ^c
30	G/G	G/A G/G	1	3591
	C/C	C/C C/C	2	3697
	C/C	C/C C/C	3	3804
	G/A	G/G G/G	4	3818
	C/C	C/C C/C	5	4123
	G/G	G/G G/G	6	4240
35	G/G	G/G G/G	7	4472
	G/G	G/G G/G	8	4499
	G/G	G/G G/G	9	4531
	G/G	G/G G/G	10	4574
	C/C	C/C C/C	11	4736
40	C/C	C/C C/C	12	4813
	C/C	C/C C/T	13	5068
	G/G	G/G G/G	14	5103
	G/G	G/G G/G	15	5150
	G/G	G/G G/A	16	5179
45	G/G	G/G G/G	17	5301
	G/G	G/G G/G	18	5333
	G/G	G/G G/G	19	5448
	G/G	G/G G/G	20	5560
	G/G	G/G G/G	21	5580
50	T/C	C/T C/C	22	5587
	G/G	G/C G/G	23	5606

^aHaplotype pairs are represented as 1st haplotype/2nd haplotype; with alleles of each haplotype shown 5' to 3' as 1st polymorphism/2nd polymorphism in each column;

55 ^bPS = polymorphic site;

^cPosition of PS in SEQ ID NO:1.

4. The method of claim 3, wherein the determining step comprises identifying the phased sequence of nucleotides present at each of PS1-PS23 on both copies of the individual's EDG6 gene.
5. A method for genotyping the endothelial differentiation, G-protein-coupled receptor 6 (EDG6) gene of an individual, comprising determining for the two copies of the EDG6 gene present in the individual the identity of the nucleotide pair at one or more polymorphic sites (PS) selected from the group consisting of PS1, PS2, PS3, PS4, PS5, PS6, PS7, PS8, PS9, PS10, PS11, PS12, PS13, PS14, PS15, PS16, PS17, PS18, PS19, PS20, PS21, PS22 and PS23, wherein the one or more PS have the location and alternative alleles shown in SEQ ID NO:1.
6. The method of claim 5, wherein the determining step comprises:
 - (a) isolating from the individual a nucleic acid mixture comprising both copies of the EDG6 gene, or a fragment thereof, that are present in the individual;
 - (b) amplifying from the nucleic acid mixture a target region containing the selected polymorphic site;
 - (c) hybridizing a primer extension oligonucleotide to one allele of the amplified target region;
 - (d) performing a nucleic acid template-dependent, primer extension reaction on the hybridized genotyping oligonucleotide in the presence of at least two different terminators of the reaction, wherein said terminators are complementary to the alternative nucleotides present at the selected polymorphic site; and
 - (e) detecting the presence and identity of the terminator in the extended genotyping oligonucleotide.
7. The method of claim 5, which comprises determining for the two copies of the EDG6 gene present in the individual the identity of the nucleotide pair at each of PS1-PS23.
8. A method for haplotyping the endothelial differentiation, G-protein-coupled receptor 6 (EDG6) gene of an individual which comprises determining, for one copy of the EDG6 gene present in the individual, the identity of the nucleotide at two or more polymorphic sites (PS) selected from the group consisting of PS1, PS2, PS3, PS4, PS5, PS6, PS7, PS8, PS9, PS10, PS11, PS12, PS13, PS14, PS15, PS16, PS17, PS18, PS19, PS20, PS21, PS22 and PS23, wherein the selected PS have the location and alternative alleles shown in SEQ ID NO:1.
9. The method of claim 8, wherein the determining step comprises:
 - (a) isolating from the individual a nucleic acid sample containing only one of the two copies of the EDG6 gene, or a fragment thereof, that is present in the individual;
 - (b) amplifying from the nucleic acid sample a target region containing the selected polymorphic site;
 - (c) hybridizing a primer extension oligonucleotide to one allele of the amplified target region;
 - (d) performing a nucleic acid template-dependent, primer extension reaction on the hybridized genotyping oligonucleotide in the presence of at least two different terminators of the reaction, wherein said terminators are complementary to the alternative nucleotides

10 present at the selected polymorphic site; and

(e) detecting the presence and identity of the terminator in the extended genotyping oligonucleotide.

10. A method for predicting a haplotype pair for the endothelial differentiation, G-protein-coupled receptor 6 (EDG6) gene of an individual comprising:

(a) identifying an EDG6 genotype for the individual, wherein the genotype comprises the nucleotide pair at two or more polymorphic sites (PS) selected from the group consisting of PS1, PS2, PS3, PS4, PS5, PS6, PS7, PS8, PS9, PS10, PS11, PS12, PS13, PS14, PS15, PS16, PS17, PS18, PS19, PS20, PS21, PS22 and PS23, wherein the selected PS have the location and alternative alleles shown in SEQ ID NO:1;

(b) enumerating all possible haplotype pairs which are consistent with the genotype;

(c) comparing the possible haplotype pairs to the haplotype pair data set forth in the table immediately below; and

(d) assigning a haplotype pair to the individual that is consistent with the data

Haplotype Pair ^a										PS ^b	PS
	18/18	17/17	5/5	16/16	17/24	5/7	17/9	17/20	Number	Position ^c	
15	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	1	3591	
	C/C	C/C	C/C	C/C	C/T	C/C	C/C	C/C	2	3697	
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	3	3804	
	G/G	G/G	A/A	G/G	G/A	A/A	G/G	G/G	4	3818	
20	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	5	4123	
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	6	4240	
	G/G	G/G	G/G	G/G	G/G	G/G	G/A	G/G	7	4472	
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	8	4499	
25	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	9	4531	
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	10	4574	
	C/C	C/C	C/C	C/C	C/C	C/T	C/C	C/C	11	4736	
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	12	4813	
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/T	13	5068	
30	G/G	G/G	G/G	G/G	G/T	G/G	G/G	G/G	14	5103	
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	15	5150	
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	16	5179	
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	17	5301	
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	18	5333	
35	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	19	5448	
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	20	5560	
	G/G	G/G	G/G	A/A	G/G	G/G	G/G	G/G	21	5580	
	T/T	C/C	C/C	C/C	C/T	C/T	C/C	C/C	22	5587	
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	23	5606	

40	Haplotype		Pair ^a						PS ^b	PS
	17/22	17/1	17/10	17/6	17/12	17/7	17/13	18/10		
45	G/G	G/A	G/G	G/G	G/G	G/G	G/G	G/G	1	3591
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	2	3697
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	3	3804
	G/G	G/G	G/G	G/A	G/G	G/A	G/G	G/G	4	3818
	C/T	C/C	C/C	C/C	C/C	C/C	C/C	C/C	5	4123
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	6	4240
50	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	7	4472
	G/G	G/G	G/A	G/G	G/G	G/G	G/G	G/A	8	4499
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	9	4531
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	10	4574
	C/C	C/C	C/C	C/C	C/T	C/C	C/C	C/C	11	4736
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	12	4813
55	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	13	5068
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	14	5103
	G/G	G/G	G/G	G/G	G/A	G/G	G/G	G/G	15	5150
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	16	5179
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	17	5301
	G/G	G/G	G/G	G/G	G/G	G/G	G/A	G/G	18	5333
60	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	19	5448
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	20	5560
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	21	5580
	C/C	C/C	C/T	C/C	C/T	C/C	T/C	T/C	22	5587
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	23	5606
65	Haplotype		Pair ^a						PS ^b	PS
	5/3	17/3	5/6	17/8	18/14	17/14	17/5	17/15		
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	1	3591
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	2	3697
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	3	3804
	A/A	G/A	A/A	G/G	G/G	G/G	G/A	G/G	4	3818
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	5	4123
	G/A	G/A	G/A	G/G	G/G	G/G	G/G	G/G	6	4240
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	7	4472
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	8	4499
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	9	4531
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	10	4574
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	11	4736
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	12	4813
80	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	13	5068
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	14	5103
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	15	5150
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	16	5179
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	17	5301
85	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	18	5333
	G/G	G/G	G/G	G/G	G/C	G/C	G/G	G/G	19	5448
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/A	20	5560
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	21	5580
	C/T	C/T	C/T	C/C	T/C	C/C	C/C	C/C	22	5587
90	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	23	5606

	Haplotype Pair ^a								PS ^b	PS
	17/23	17/11	5/4	18/6	17/18	17/16	18/3	17/21	Number	Position ^c
95	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	1	3591
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	2	3697
	C/T	C/C	C/C	C/C	C/C	C/C	C/C	C/C	3	3804
	G/G	G/G	A/A	G/A	G/G	G/G	G/A	G/G	4	3818
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	5	4123
	G/G	G/G	G/G	G/G	G/G	G/G	G/A	G/G	6	4240
100	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	7	4472
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	8	4499
	G/G	G/A	G/G	G/G	G/G	G/G	G/G	G/G	9	4531
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/T	10	4574
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	11	4736
	C/C	C/T	C/C	C/C	C/C	C/C	C/C	C/C	12	4813
105	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	13	5068
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	14	5103
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	15	5150
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	16	5179
	G/G	G/G	G/A	G/G	G/G	G/G	G/G	G/G	17	5301
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	18	5333
110	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	19	5448
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	20	5560
	G/G	G/G	G/G	G/G	G/G	G/A	G/G	G/G	21	5580
	C/T	C/T	C/C	T/T	C/T	C/C	T/T	C/C	22	5587
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	23	5606

	Haplotype	Pair ^a	PS ^b	PS	
	18/5	17/2	17/19	Number	Position ^c
120	G/G	G/A	G/G	1	3591
	C/C	C/C	C/C	2	3697
	C/C	C/C	C/C	3	3804
	G/A	G/G	G/G	4	3818
	C/C	C/C	C/C	5	4123
	G/G	G/G	G/G	6	4240
125	G/G	G/G	G/G	7	4472
	G/G	G/G	G/G	8	4499
	G/G	G/G	G/G	9	4531
	G/G	G/G	G/G	10	4574
	C/C	C/C	C/C	11	4736
	C/C	C/C	C/C	12	4813
130	C/C	C/C	C/T	13	5068
	G/G	G/G	G/G	14	5103
	G/G	G/G	G/G	15	5150
	G/G	G/G	G/A	16	5179
	G/G	G/G	G/G	17	5301
	G/G	G/G	G/G	18	5333
135	G/G	G/G	G/G	19	5448
	G/G	G/G	G/G	20	5560
	G/G	G/G	G/G	21	5580
	T/C	C/T	C/C	22	5587
	G/G	G/C	G/G	23	5606

145 ^aHaplotype pairs are represented as 1st haplotype/2nd haplotype; with alleles of each haplotype shown 5' to 3' as 1st polymorphism/2nd polymorphism in each column;

^bPS = polymorphic site;

^cPosition of PS in SEQ ID NO:1.

11. The method of claim 10, wherein the identified genotype of the individual comprises the nucleotide pair at each of PS1-PS23, which have the location and alternative alleles shown in SEQ ID NO:1.

12. A method for identifying an association between a trait and at least one haplotype or haplotype pair of the endothelial differentiation, G-protein-coupled receptor 6 (EDG6) gene which comprises comparing the frequency of the haplotype or haplotype pair in a population exhibiting the trait with the frequency of the haplotype or haplotype pair in a reference population, wherein the haplotype is selected from haplotypes 1-24 shown in the table presented immediately below, wherein each of the haplotypes comprises a set of polymorphisms whose locations and identities are set forth in the table immediately below:

10

Haplotype Number ^a											PS ^b	PS
1	2	3	4	5	6	7	8	9	10	Number	Position ^c	
A	A	G	G	G	G	G	G	G	G	1	3591	
15	C	C	C	C	C	C	C	C	C	2	3697	
C	C	C	C	C	C	C	C	C	C	3	3804	
G	G	A	A	A	A	A	G	G	G	4	3818	
C	C	C	C	C	C	C	C	C	C	5	4123	
G	G	A	G	G	G	G	A	G	G	6	4240	
20	G	G	G	G	G	G	G	A	G	7	4472	
G	G	G	G	G	G	G	G	G	A	8	4499	
G	G	G	G	G	G	G	G	G	G	9	4531	
G	G	G	G	G	G	G	G	G	G	10	4574	
25	C	C	C	C	C	T	C	C	C	11	4736	
C	C	C	C	C	C	C	C	C	C	12	4813	
C	C	C	C	C	C	C	C	C	C	13	5068	
G	G	G	G	G	G	G	G	G	G	14	5103	
G	G	G	G	G	G	G	G	G	G	15	5150	
G	G	G	G	G	G	G	G	G	G	16	5179	
30	G	G	G	A	G	G	G	G	G	17	5301	
G	G	G	G	G	G	G	G	G	G	18	5333	
G	G	G	G	G	G	G	G	G	G	19	5448	
G	G	G	G	G	G	G	G	G	G	20	5560	
G	G	G	G	G	G	G	G	G	G	21	5580	
35	C	T	T	C	C	T	T	C	T	C	22	5587
G	C	G	G	G	G	G	G	G	G	23	5606	

Haplotype Number ^a												PS ^b	PS
	11	12	13	14	15	16	17	18	19	20	Number	Position ^c	
40	G	G	G	G	G	G	G	G	G	G	1	3591	
	C	C	C	C	C	C	C	C	C	C	2	3697	
	C	C	C	C	C	C	C	C	C	C	3	3804	
	G	G	G	G	G	G	G	G	G	G	4	3818	
45	C	C	C	C	C	C	C	C	C	C	5	4123	
	G	G	G	G	G	G	G	G	G	G	6	4240	
	G	G	G	G	G	G	G	G	G	G	7	4472	
	G	G	G	G	G	G	G	G	G	G	8	4499	
50	A	G	G	G	G	G	G	G	G	G	9	4531	
	G	G	G	G	G	G	G	G	G	G	10	4574	
	C	C	C	C	C	C	C	C	C	C	11	4736	
	T	C	C	C	C	C	C	C	C	C	12	4813	
55	C	C	C	C	C	C	C	T	T	T	13	5068	
	G	G	G	G	G	G	G	G	G	G	14	5103	
	G	A	G	G	G	G	G	G	G	G	15	5150	
	G	G	G	G	G	G	G	A	G	G	16	5179	
60	G	G	G	G	G	G	G	G	G	G	17	5301	
	G	G	A	G	G	G	G	G	G	G	18	5333	
	G	G	G	C	G	G	G	G	G	G	19	5448	
	G	G	G	G	A	G	G	G	G	G	20	5560	
65	G	G	G	G	A	G	G	G	G	G	21	5580	
	T	C	C	C	C	C	T	C	C	C	22	5587	
	G	G	G	G	G	G	G	G	G	G	23	5606	

Haplotype Number ^a					PS ^b	PS
	21	22	23	24	Number	Position ^c
65	G	G	G	G	1	3591
	C	C	C	T	2	3697
	C	C	T	C	3	3804
	G	G	G	A	4	3818
70	C	T	C	C	5	4123
	G	G	G	G	6	4240
	G	G	G	G	7	4472
	G	G	G	G	8	4499
75	G	G	G	G	9	4531
	T	G	G	G	10	4574
	C	C	C	C	11	4736
	C	C	C	C	12	4813
80	C	C	C	C	13	5068
	G	G	G	T	14	5103
	G	G	G	G	15	5150
	G	G	G	G	16	5179
85	G	G	G	G	17	5301
	G	G	G	G	18	5333
	G	G	G	G	19	5448
	G	G	G	G	20	5560
90	G	G	G	G	21	5580
	C	C	T	C	22	5587
	G	G	G	G	23	5606

^aAlleles for haplotypes are presented 5' to 3' in each column

^bPS = polymorphic site;

^cPosition of PS in SEQ ID NO:1;

and wherein the haplotype pair is selected from the haplotype pairs shown in the table

immediately below, wherein each of the EDG6 haplotype pairs consists of first and second
 95 haplotypes which comprise first and second sets of polymorphisms whose locations and
 identities are set forth in the table immediately below:

	Haplotype Pair ^a								PS ^b	PS
	18/18	17/17	5/5	16/16	17/24	5/7	17/9	17/20	Number	Position ^c
100	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	1	3591
	C/C	C/C	C/C	C/C	C/T	C/C	C/C	C/C	2	3697
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	3	3804
	G/G	G/G	A/A	G/G	G/A	A/A	G/G	G/G	4	3818
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	5	4123
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	6	4240
105	G/G	G/G	G/G	G/G	G/G	G/G	G/A	G/G	7	4472
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	8	4499
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	9	4531
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	10	4574
	C/C	C/C	C/C	C/C	C/C	C/T	C/C	C/C	11	4736
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	12	4813
110	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	13	5068
	G/G	G/G	G/G	G/G	G/T	G/G	G/G	G/G	14	5103
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	15	5150
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	16	5179
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	17	5301
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	18	5333
115	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	19	5448
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	20	5560
	G/G	G/G	G/G	A/A	G/G	G/G	G/G	G/G	21	5580
	T/T	C/C	C/C	C/C	C/C	C/T	C/T	C/C	22	5587
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	23	5606
	Haplotype Pair ^a								PS ^b	PS
125	17/22	17/1	17/10	17/6	17/12	17/7	17/13	18/10	Number	Position ^c
	G/G	G/A	G/G	G/G	G/G	G/G	G/G	G/G	1	3591
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	2	3697
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	3	3804
	G/G	G/G	G/A	G/G	G/A	G/A	G/G	G/G	4	3818
	C/T	C/C	C/C	C/C	C/C	C/C	C/C	C/C	5	4123
130	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	6	4240
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	7	4472
	G/G	G/A	G/G	G/G	G/G	G/G	G/A	G/A	8	4499
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	9	4531
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	10	4574
	C/C	C/C	C/C	C/C	C/C	C/T	C/C	C/C	11	4736
135	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	12	4813
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	13	5068
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	14	5103
	G/G	G/G	G/G	G/G	G/A	G/G	G/G	G/G	15	5150
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	16	5179
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	17	5301
140	G/G	G/G	G/G	G/G	G/G	G/G	G/A	G/G	18	5333
	G/G	G/G	G/G	G/G	G/G	G/G	G/A	G/G	19	5448
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	20	5560
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	21	5580
	C/C	C/C	C/C	C/T	C/C	C/T	C/C	T/C	22	5587
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	23	5606

		Haplotype	Pair ^a							PS ^b	PS
150		5/3	17/3	5/6	17/8	18/14	17/14	17/5	17/15	Number	Position ^c
		G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	1	3591
		C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	2	3697
		C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	3	3804
		A/A	G/A	A/A	G/G	G/G	G/G	G/A	G/G	4	3818
155		C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	5	4123
		G/A	G/A	G/G	G/A	G/G	G/G	G/G	G/G	6	4240
		G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	7	4472
		G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	8	4499
		G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	9	4531
160		G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	10	4574
		C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	11	4736
		C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	12	4813
		C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	13	5068
		G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	14	5103
165		G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	15	5150
		G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	16	5179
		G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	17	5301
		G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	18	5333
		G/G	G/G	G/G	G/C	G/C	G/C	G/G	G/G	19	5448
170		G/G	G/G	G/G	G/G	G/G	G/G	G/A	G/A	20	5560
		G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	21	5580
		C/T	C/T	C/C	T/C	C/C	C/C	C/C	C/C	22	5587
		G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	23	5606
175		Haplotype	Pair ^a							PS ^b	PS
		17/23	17/11	5/4	18/6	17/18	17/16	18/3	17/21	Number	Position ^c
		G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	1	3591
		C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	2	3697
		C/T	C/C	C/C	C/C	C/C	C/C	C/C	C/C	3	3804
180		G/G	G/G	A/A	G/A	G/G	G/G	G/A	G/G	4	3818
		C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	5	4123
		G/G	G/G	G/G	G/G	G/G	G/G	G/A	G/G	6	4240
		G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	7	4472
		G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	8	4499
185		G/G	G/A	G/G	G/G	G/G	G/G	G/G	G/G	9	4531
		G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/T	10	4574
		C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	11	4736
		C/C	C/T	C/C	C/C	C/C	C/C	C/C	C/C	12	4813
		C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	13	5068
190		G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	14	5103
		G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	15	5150
		G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	16	5179
		G/G	G/G	G/A	G/G	G/G	G/G	G/G	G/G	17	5301
		G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	18	5333
195		G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	19	5448
		G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	20	5560
		G/G	G/G	G/G	G/G	G/A	G/A	G/G	G/G	21	5580
		C/T	C/T	C/C	T/T	C/T	C/C	T/T	C/C	22	5587
		G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	23	5606
200											

	Haplotype	Pair ^a	PS ^b	PS	
	18/5	17/2	17/19	Number	Position ^c
205	G/G	G/A	G/G	1	3591
	C/C	C/C	C/C	2	3697
	C/C	C/C	C/C	3	3804
	G/A	G/G	G/G	4	3818
	C/C	C/C	C/G	5	4123
	G/G	G/G	G/G	6	4240
	G/G	G/G	G/G	7	4472
210	G/G	G/G	G/G	8	4499
	G/G	G/G	G/G	9	4531
	G/G	G/G	G/G	10	4574
	C/C	C/C	C/C	11	4736
	C/C	C/C	C/C	12	4813
215	C/C	C/C	C/T	13	5068
	G/G	G/G	G/G	14	5103
	G/G	G/G	G/G	15	5150
	G/G	G/G	G/A	16	5179
	G/G	G/G	G/G	17	5301
220	G/G	G/G	G/G	18	5333
	G/G	G/G	G/G	19	5448
	G/G	G/G	G/G	20	5560
	G/G	G/G	G/G	21	5580
	T/C	C/T	C/C	22	5587
225	G/G	G/C	G/G	23	5606

^aHaplotype pairs are represented as 1st haplotype/2nd haplotype; with alleles of each haplotype shown 5' to 3' as 1st polymorphism/2nd polymorphism in each column;

^bPS = polymorphic site;

^cPosition of PS in SEQ ID NO:1;

wherein a higher frequency of the haplotype or haplotype pair in the trait population than in the reference population indicates the trait is associated with the haplotype or haplotype pair.

13. The method of claim 12, wherein the trait is a clinical response to a drug targeting EDG6.
14. An isolated genotyping oligonucleotide for detecting a polymorphism in the endothelial differentiation, G-protein-coupled receptor 6 (EDG6) gene at a polymorphic site (PS) selected from the group consisting of PS1, PS2, PS3, PS4, PS5, PS6, PS7, PS8, PS9, PS10, PS11, PS12, PS13, PS14, PS15, PS16, PS17, PS18, PS19, PS20, PS21, PS22 and PS23, wherein the selected PS have the location and alternative alleles shown in SEQ ID NO:1.
15. The isolated genotyping oligonucleotide of claim 14, which is an allele-specific oligonucleotide that specifically hybridizes to an allele of the EDG6 gene at a region containing the polymorphic site.
16. The allele-specific oligonucleotide of claim 15, which comprises a nucleotide sequence selected from the group consisting of SEQ ID NOS:4-26, the complements of SEQ ID NOS:4-26, and SEQ ID NOS:27-72.
17. The isolated genotyping oligonucleotide of claim 14, which is a primer-extension oligonucleotide.
18. The primer-extension oligonucleotide of claim 17, which comprises a nucleotide sequence

selected from the group consisting of SEQ ID NOS:73-118.

19. A kit for genotyping the endothelial differentiation, G-protein-coupled receptor 6 (EDG6) gene of an individual, which comprises a set of oligonucleotides designed to genotype each of polymorphic sites (PS) PS1, PS2, PS3, PS4, PS5, PS6, PS7, PS8, PS9, PS10, PS11, PS12, PS13, PS14, PS15, PS16, PS17, PS18, PS19, PS20, PS21, PS22 and PS23, wherein the selected PS have the location and alternative alleles shown in SEQ ID NO:1.

20. An isolated polynucleotide comprising a nucleotide sequence selected from the group consisting of:

(a) a first nucleotide sequence which comprises a endothelial differentiation, G-protein-coupled receptor 6 (EDG6) isogene, wherein the EDG6 isogene is selected from the group consisting of isogenes 1- 4 and 6 - 24 shown in the table immediately below and wherein each of the isogenes comprises the regions of SEQ ID NO:1 shown in the table immediately below and wherein each of the isogenes 1- 4 and 6 - 24 is further defined by the corresponding set of polymorphisms whose locations and identities are set forth in the table immediately below

1	2	3	4	5	6	7	8	9	10	No.	PS ^b	PS	SEQ ID	Region
											Pos. ^c	No.	Examined ^d	
A	A	G	G	G	G	G	G	G	G	1	3591	1	3484-5771	
C	C	C	C	C	C	C	C	C	C	2	3697	1	3484-5771	
C	C	C	C	C	C	C	C	C	C	3	3804	1	3484-5771	
G	G	A	A	A	A	A	G	G	G	4	3818	1	3484-5771	
C	C	C	C	C	C	C	C	C	C	5	4123	1	3484-5771	
G	G	A	G	G	G	G	A	G	G	6	4240	1	3484-5771	
G	G	G	G	G	G	G	G	A	G	7	4472	1	3484-5771	
G	G	G	G	G	G	G	G	G	A	8	4499	1	3484-5771	
G	G	G	G	G	G	G	G	G	G	9	4531	1	3484-5771	
G	G	G	G	G	G	G	G	G	G	10	4574	1	3484-5771	
C	C	C	C	C	C	T	C	C	C	11	4736	1	3484-5771	
C	C	C	C	C	C	C	C	C	C	12	4813	1	3484-5771	
C	C	C	C	C	C	C	C	C	C	13	5068	1	3484-5771	
G	G	G	G	G	G	G	G	G	G	14	5103	1	3484-5771	
G	G	G	G	G	G	G	G	G	G	15	5150	1	3484-5771	
G	G	G	G	G	G	G	G	G	G	16	5179	1	3484-5771	
G	G	G	A	G	G	G	G	G	G	17	5301	1	3484-5771	
G	G	G	G	G	G	G	G	G	G	18	5333	1	3484-5771	
G	G	G	G	G	G	G	G	G	G	19	5448	1	3484-5771	
G	G	G	G	G	G	G	G	G	G	20	5560	1	3484-5771	
G	G	G	G	G	G	G	G	G	G	21	5580	1	3484-5771	
C	T	T	C	C	T	T	C	T	C	22	5587	1	3484-5771	
G	C	G	G	G	G	G	G	G	G	23	5606	1	3484-5771	

Isogene Number ^a												PS ^b	PS	SEQ	ID	Region
11	12	13	14	15	16	17	18	19	20	No.	Pos. ^c	No.	Examined ^d			
G	G	G	G	G	G	G	G	G	G	1	3591	1	3484-5771			
C	C	C	C	C	C	C	C	C	C	2	3697	1	3484-5771			
C	C	C	C	C	C	C	C	C	C	3	3804	1	3484-5771			
G	G	G	G	G	G	G	G	G	G	4	3818	1	3484-5771			
C	C	C	C	C	C	C	C	C	C	5	4123	1	3484-5771			
G	G	G	G	G	G	G	G	G	G	6	4240	1	3484-5771			
G	G	G	G	G	G	G	G	G	G	7	4472	1	3484-5771			
G	G	G	G	G	G	G	G	G	G	8	4499	1	3484-5771			
A	G	G	G	G	G	G	G	G	G	9	4531	1	3484-5771			
G	G	G	G	G	G	G	G	G	G	10	4574	1	3484-5771			
C	C	C	C	C	C	C	C	C	C	11	4736	1	3484-5771			
T	C	C	C	C	C	C	C	C	C	12	4813	1	3484-5771			
C	C	C	C	C	C	C	C	T	T	13	5068	1	3484-5771			
G	G	G	G	G	G	G	G	G	G	14	5103	1	3484-5771			
G	A	G	G	G	G	G	G	G	G	15	5150	1	3484-5771			
G	G	G	G	G	G	G	G	A	G	16	5179	1	3484-5771			
G	G	G	G	G	G	G	G	G	G	17	5301	1	3484-5771			
G	G	A	G	G	G	G	G	G	G	18	5333	1	3484-5771			
G	G	G	C	G	G	G	G	G	G	19	5448	1	3484-5771			
G	G	G	G	A	G	G	G	G	G	20	5560	1	3484-5771			
G	G	G	G	G	A	G	G	G	G	21	5580	1	3484-5771			
T	C	C	C	C	C	C	T	C	C	22	5587	1	3484-5771			
G	G	G	G	G	G	G	G	G	G	23	5606	1	3484-5771			

Isogene Number ^a				PS ^b	PS	SEQ	ID	Region
21	22	23	24	No.	Pos. ^c	No.	Examined ^d	
G	G	G	G	1	3591	1	3484-5771	
C	C	C	T	2	3697	1	3484-5771	
C	C	T	C	3	3804	1	3484-5771	
G	G	G	A	4	3818	1	3484-5771	
C	T	C	C	5	4123	1	3484-5771	
G	G	G	G	6	4240	1	3484-5771	
G	G	G	G	7	4472	1	3484-5771	
G	G	G	G	8	4499	1	3484-5771	
G	G	G	G	9	4531	1	3484-5771	
T	G	G	G	10	4574	1	3484-5771	
C	C	C	C	11	4736	1	3484-5771	
C	C	C	C	12	4813	1	3484-5771	
C	C	C	C	13	5068	1	3484-5771	
G	G	G	T	14	5103	1	3484-5771	
G	G	G	G	15	5150	1	3484-5771	
G	G	G	G	16	5179	1	3484-5771	
G	G	G	G	17	5301	1	3484-5771	
G	G	G	G	18	5333	1	3484-5771	
G	G	G	G	19	5448	1	3484-5771	
G	G	G	G	20	5560	1	3484-5771	
G	G	G	G	21	5580	1	3484-5771	
C	C	T	C	22	5587	1	3484-5771	
G	G	G	G	23	5606	1	3484-5771	

^aAlleles for isogenes are presented 5' to 3' in each column

^bPS = polymorphic site;

^cPosition of PS in SEQ ID NO:1;

^dRegion examined represents the nucleotide positions defining the start and stop positions within the SEQ ID NO of the sequenced region.

(b) a second nucleotide sequence which comprises a fragment of the first nucleotide sequence, wherein the fragment comprises one or more polymorphisms selected from the group consisting of adenine at PS1, thymine at PS2, thymine at PS3, guanine at PS4, thymine at PS5, adenine at PS6, adenine at PS7, adenine at PS8, adenine at PS9, thymine at PS10, thymine at PS11, thymine at PS12, thymine at PS13, thymine at PS14, adenine at PS15, adenine at PS16, adenine at PS17, adenine at PS18, cytosine at PS19, adenine at PS20, adenine at PS21, thymine at PS22 and cytosine at PS23, wherein the selected polymorphism has the location set forth in the table immediately above; and

(c) a third nucleotide sequence which is complementary to the first or second nucleotide sequence.

21. The isolated polynucleotide of claim 20, which is a DNA molecule and comprises both the first and third nucleotide sequences and further comprises expression regulatory elements operably linked to the first nucleotide sequence.

22. A recombinant nonhuman organism transformed or transfected with the isolated polynucleotide of claim 20, wherein the organism expresses an EDG6 protein encoded by the first nucleotide sequence.

23. The recombinant nonhuman organism of claim 22, which is a transgenic animal.

24. The isolated polynucleotide of claim 20 which consists of the second nucleotide sequence.

25. An isolated polynucleotide comprising a nucleotide sequence selected from the group consisting of:

(a) a coding sequence for a endothelial differentiation, G-protein-coupled receptor 6 (EDG6) isogene wherein the coding sequence is defined by a haplotype selected from the group consisting of 3c, 7c-12c, 19c-22c, and 24c shown in the table immediately below and wherein the coding sequence comprises SEQ ID NO:2 except at each of the polymorphic sites which have the locations and polymorphisms set forth in the table immediately below:

Coding	Sequence	Haplotype Number ^a												PS ^b No.	PS Position ^c
		3c	7c	8c	9c	10c	11c	12c	19c	20c	21c	22c	24c		
C	C	C	C	C	C	C	C	C	C	C	T	C	5	114	
A	G	A	G	G	G	G	G	G	G	G	G	G	6	231	
G	G	G	A	G	G	G	G	G	G	G	G	G	7	463	
G	G	G	G	A	G	G	G	G	G	G	G	G	8	490	
G	G	G	G	G	A	G	G	G	G	G	G	G	9	522	
G	G	G	G	G	G	G	G	G	T	G	G	G	10	565	
C	T	C	C	C	C	C	C	C	C	C	C	C	11	727	
C	C	C	C	C	T	C	C	C	C	C	C	C	12	804	
C	C	C	C	C	C	C	T	T	C	C	C	C	13	1059	
G	G	G	G	G	G	G	G	G	G	G	T	G	14	1094	
G	G	G	G	G	G	A	G	G	G	G	G	G	15	1141	

^aAlleles for coding sequence haplotypes are presented 5' to 3' in each column; the numerical portion of the coding sequence haplotype number represents the number of the

parent full EDG6 haplotype;

^bPS = polymorphic site;

^cPosition of PS in SEQ ID NO:2;

and

(b) a fragment of the coding sequence, wherein the fragment comprises at least one polymorphism selected from the group consisting of thymine at a position corresponding to nucleotide 114, adenine at a position corresponding to nucleotide 231, adenine at a position corresponding to nucleotide 463, adenine at a position corresponding to nucleotide 490, adenine at a position corresponding to nucleotide 522, thymine at a position corresponding to nucleotide 565, thymine at a position corresponding to nucleotide 727, thymine at a position corresponding to nucleotide 804, thymine at a position corresponding to nucleotide 1059, thymine at a position corresponding to nucleotide 1094 and adenine at a position corresponding to nucleotide 1141, wherein said positions in the coding sequence and the fragment refer to SEQ ID NO:2.

26. A recombinant nonhuman organism transformed or transfected with the isolated polynucleotide of claim 25, wherein the organism expresses a endothelial differentiation, G-protein-coupled receptor 6 (EDG6) protein encoded by the polymorphic variant sequence.

27. The recombinant nonhuman organism of claim 26, which is a transgenic animal.

28. An isolated polypeptide comprising an amino acid sequence which is a polymorphic variant of a reference sequence for the endothelial differentiation, G-protein-coupled receptor 6 (EDG6) protein or a fragment thereof, wherein the reference sequence comprises SEQ ID NO:3 and the polymorphic variant comprises one or more variant amino acids selected from the group consisting of arginine at a position corresponding to amino acid position 155, serine at a position corresponding to amino acid position 164, serine at a position corresponding to amino acid position 189, cysteine at a position corresponding to amino acid position 243, leucine at a position corresponding to amino acid position 365 and methionine at a position corresponding to amino acid position 381.

29. An isolated monoclonal antibody specific for and immunoreactive with the isolated polypeptide of claim 28.

30. A method for screening for drugs targeting the isolated polypeptide of claim 28 which comprises contacting the EDG6 polymorphic variant with a candidate agent and assaying for binding activity.

31. A computer system for storing and analyzing polymorphism data for the endothelial differentiation, G-protein-coupled receptor 6 gene, comprising:

- a central processing unit (CPU);
- a communication interface;
- a display device;
- an input device; and

(e) a database containing the polymorphism data;
 wherein the polymorphism data comprises the haplotypes set forth in the table immediately below:

Haplotype Number ^a											PS ^b	PS
	1	2	3	4	5	6	7	8	9	10	Number	Position ^c
10	A	A	G	G	G	G	G	G	G	G	1	3591
	C	C	C	C	C	C	C	C	C	C	2	3697
	C	C	C	C	C	C	C	C	C	C	3	3804
15	G	G	A	A	A	A	A	G	G	G	4	3818
	C	C	C	C	C	C	C	C	C	C	5	4123
	G	G	A	G	G	G	G	A	G	G	6	4240
	G	G	G	G	G	G	G	G	A	G	7	4472
	G	G	G	G	G	G	G	G	G	A	8	4499
20	G	G	G	G	G	G	G	G	G	G	9	4531
	G	G	G	G	G	G	G	G	G	G	10	4574
	C	C	C	C	C	C	T	C	C	C	11	4736
	C	C	C	C	C	C	C	C	C	C	12	4813
25	C	C	C	C	C	C	C	C	C	C	13	5068
	G	G	G	G	G	G	G	G	G	G	14	5103
	G	G	G	G	G	G	G	G	G	G	15	5150
	G	G	G	G	G	G	G	G	G	G	16	5179
	G	G	G	A	G	G	G	G	G	G	17	5301
30	G	G	G	G	G	G	G	G	G	G	18	5333
	G	G	G	G	G	G	G	G	G	G	19	5448
	G	G	G	G	G	G	G	G	G	G	20	5560
	G	G	G	G	G	G	G	G	G	G	21	5580
	C	T	T	C	C	T	T	C	T	C	22	5587
35	G	C	G	G	G	G	G	G	G	G	23	5606

Haplotype Number ^a												PS ^b	PS
												Number	Position ^c
11	12	13	14	15	16	17	18	19	20			1	3591
G	G	G	G	G	G	G	G	G	G			2	3697
40	C	C	C	C	C	C	C	C	C			3	3804
G	G	G	G	G	G	G	G	G	G			4	3818
C	C	C	C	C	C	C	C	C	C			5	4123
G	G	G	G	G	G	G	G	G	G			6	4240
45	G	G	G	G	G	G	G	G	G			7	4472
G	G	G	G	G	G	G	G	G	G			8	4499
A	G	G	G	G	G	G	G	G	G			9	4531
G	G	G	G	G	G	G	G	G	G			10	4574
C	C	C	C	C	C	C	C	C	C			11	4736
50	T	C	C	C	C	C	C	C	C			12	4813
C	C	C	C	C	C	C	C	T	T			13	5068
G	G	G	G	G	G	G	G	G	G			14	5103
G	A	G	G	G	G	G	G	G	G			15	5150
G	G	G	G	G	G	G	G	A	G			16	5179
55	G	G	G	G	G	G	G	G	G			17	5301
G	G	A	G	G	G	G	G	G	G			18	5333
G	G	G	C	G	G	G	G	G	G			19	5448
G	G	G	G	A	G	G	G	G	G			20	5560
G	G	G	G	G	A	G	G	G	G			21	5580
60	T	C	C	C	C	C	T	C	C			22	5587
G	G	G	G	G	G	G	G	G	G			23	5606

Haplotype Number ^a				PS ^b	PS
21	22	23	24	Number	Position ^c
G	G	G	G	1	3591
65	C	C	C	2	3697
C	C	T	C	3	3804
G	G	G	A	4	3818
C	T	C	C	5	4123
G	G	G	G	6	4240
70	G	G	G	7	4472
G	G	G	G	8	4499
G	G	G	G	9	4531
T	G	G	G	10	4574
C	C	C	C	11	4736
75	C	C	C	12	4813
C	C	C	C	13	5068
G	G	G	T	14	5103
G	G	G	G	15	5150
80	G	G	G	16	5179
G	G	G	G	17	5301
G	G	G	G	18	5333
G	G	G	G	19	5448
G	G	G	G	20	5560
85	G	G	G	21	5580
C	C	T	C	22	5587
G	G	G	G	23	5606

^aAlleles for haplotypes are presented 5' to 3' in each column

^bPS = polymorphic site;

^cPosition of PS in SEQ ID NO:1;

	Haplotype Pair ^a								PS ^b	PS	
	18/18	17/17	5/5	16/16	17/24	5/7	17/9	17/20	Number	Position ^c	
95	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	1	3591	
	C/C	C/C	C/C	C/C	C/T	C/C	C/C	C/C	2	3697	
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	3	3804	
	G/G	G/G	A/A	G/G	G/A	A/A	G/G	G/G	4	3818	
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	5	4123	
100	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	6	4240	
	G/G	G/G	G/G	G/G	G/G	G/G	G/A	G/G	7	4472	
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	8	4499	
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	9	4531	
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	10	4574	
105	C/C	C/C	C/C	C/C	C/C	C/T	C/C	C/C	11	4736	
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	12	4813	
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/T	13	5068	
	G/G	G/G	G/G	G/G	G/T	G/G	G/G	G/G	14	5103	
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	15	5150	
110	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	16	5179	
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	17	5301	
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	18	5333	
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	19	5448	
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	20	5560	
115	G/G	G/G	G/G	A/A	G/G	G/G	G/G	G/G	21	5580	
	T/T	C/C	C/C	C/C	C/T	C/T	C/C	C/C	22	5587	
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	23	5606	
	Haplotype Pair ^a								PS ^b	PS	
	120	17/22	17/1	17/10	17/6	17/12	17/7	17/13	18/10	Number	Position ^c
120	G/G	G/A	G/G	G/G	G/G	G/G	G/G	G/G	G/G	1	3591
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	2	3697
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	3	3804
	G/G	G/G	G/G	G/A	G/G	G/A	G/G	G/G	G/G	4	3818
	C/T	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	5	4123
125	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	6	4240
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	7	4472
	G/G	G/G	G/A	G/G	G/G	G/G	G/G	G/A	G/A	8	4499
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	9	4531
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	10	4574
130	C/C	C/C	C/C	C/C	C/C	C/T	C/C	C/C	C/C	11	4736
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	12	4813
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	13	5068
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	14	5103
	G/G	G/G	G/G	G/G	G/A	G/G	G/G	G/G	G/G	15	5150
135	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	16	5179
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	17	5301
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	18	5333
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	19	5448
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	20	5560
140	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	21	5580
	C/C	C/C	C/C	C/T	C/C	C/T	C/C	T/C	T/C	22	5587
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	23	5606

145	Haplotype	Pair ^a								PS ^b	PS
		5/3	17/3	5/6	17/8	18/14	17/14	17/5	17/15		
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	1	3591
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	2	3697
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	3	3804
150	A/A	G/A	A/A	G/G	G/G	G/G	G/A	G/G	G/G	4	3818
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	5	4123
	G/A	G/A	G/G	G/A	G/G	G/G	G/G	G/G	G/G	6	4240
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	7	4472
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	8	4499
155	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	9	4531
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	10	4574
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	11	4736
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	12	4813
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	13	5068
160	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	14	5103
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	15	5150
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	16	5179
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	17	5301
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	18	5333
165	G/G	G/G	G/G	G/G	G/C	G/C	G/G	G/G	G/G	19	5448
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/A	G/A	20	5560
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	21	5580
	C/T	C/T	C/C	T/C	C/C	C/C	C/C	C/C	C/C	22	5587
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	23	5606

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	Haplotype Pair ^a							PS ^b	PS	
	17/23	17/11	5/4	18/6	17/18	17/16	18/3	17/21	Number	Position ^c
175	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	1	3591
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	2	3697
	C/T	C/C	C/C	C/C	C/C	C/C	C/C	C/C	3	3804
	G/G	G/G	A/A	G/A	G/G	G/G	G/A	G/G	4	3818
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	5	4123
	G/G	G/G	G/G	G/G	G/G	G/G	G/A	G/G	6	4240
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	7	4472
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	8	4499
180	G/G	G/A	G/G	G/G	G/G	G/G	G/G	G/G	9	4531
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/T	10	4574
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	11	4736
	C/C	C/T	C/C	C/C	C/C	C/C	C/C	C/C	12	4813
	C/C	C/C	C/C	C/C	C/C	C/C	C/C	C/C	13	5068
185	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	14	5103
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	15	5150
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	16	5179
	G/G	G/G	G/A	G/G	G/G	G/G	G/G	G/G	17	5301
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	18	5333
190	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	19	5448
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	20	5560
	G/G	G/G	G/G	G/G	G/G	G/A	G/G	G/G	21	5580
	C/T	C/T	C/C	T/T	C/T	C/C	T/T	C/C	22	5587
	G/G	G/G	G/G	G/G	G/G	G/G	G/G	G/G	23	5606

	Haplotype	Pair ^a	PS ^b	PS	
	18/5	17/2	17/19	Number	Position ^c
200	G/G	G/A	G/G	1	3591
	C/C	C/C	C/C	2	3697
	C/C	C/C	C/C	3	3804
	G/A	G/G	G/G	4	3818
	C/C	C/C	C/C	5	4123
	G/G	G/G	G/G	6	4240
	G/G	G/G	G/G	7	4472
	G/G	G/G	G/G	8	4499
205	G/G	G/G	G/G	9	4531
	G/G	G/G	G/G	10	4574
	C/C	C/C	C/C	11	4736
	C/C	C/C	C/C	12	4813
	C/C	C/C	C/T	13	5068
210	G/G	G/G	G/G	14	5103
	G/G	G/G	G/G	15	5150
	G/G	G/G	G/A	16	5179
	G/G	G/G	G/G	17	5301
	G/G	G/G	G/G	18	5333
215	G/G	G/G	G/G	19	5448
	G/G	G/G	G/G	20	5560
	G/G	G/G	G/G	21	5580
	T/C	C/T	C/C	22	5587
	G/G	G/C	G/G	23	5606

^aHaplotype pairs are represented as 1st Haplotype/2nd Haplotype; with alleles of each haplotype shown 5' to 3' as 1st polymorphism/2nd polymorphism in each column;

^bPS = polymorphic site

^cLocation of PS in SEQ ID NO:1.

32. A genome anthology for the endothelial differentiation, G-protein-coupled receptor 6 (EDG6) gene which comprises EDG6 isogenes defined by any one of haplotypes 1-24 set forth in the table shown below:

5	Haplotype Number ^a										PS ^b Number	PS Position ^c
	1	2	3	4	5	6	7	8	9	10		
	A	A	G	G	G	G	G	G	G	G	1	3591
	C	C	C	C	C	C	C	C	C	C	2	3697
	C	C	C	C	C	C	C	C	C	C	3	3804
10	G	G	A	A	A	A	A	G	G	G	4	3818
	C	C	C	C	C	C	C	C	C	C	5	4123
	G	G	A	G	G	G	G	A	G	G	6	4240
	G	G	G	G	G	G	G	G	A	G	7	4472
	G	G	G	G	G	G	G	G	G	A	8	4499
15	G	G	G	G	G	G	G	G	G	G	9	4531
	G	G	G	G	G	G	G	G	G	G	10	4574
	C	C	C	C	C	C	T	C	C	C	11	4736
	C	C	C	C	C	C	C	C	C	C	12	4813
	C	C	C	C	C	C	C	C	C	C	13	5068
20	G	G	G	G	G	G	G	G	G	G	14	5103
	G	G	G	G	G	G	G	G	G	G	15	5150
	G	G	G	G	G	G	G	G	G	G	16	5179
	G	G	G	A	G	G	G	G	G	G	17	5301
	G	G	G	G	G	G	G	G	G	G	18	5333
25	G	G	G	G	G	G	G	G	G	G	19	5448
	G	G	G	G	G	G	G	G	G	G	20	5560
	G	G	G	G	G	G	G	G	G	G	21	5580
	C	T	T	C	C	T	T	C	T	C	22	5587
	G	C	G	G	G	G	G	G	G	G	23	5606
30	Haplotype Number ^a										PS ^b Number	PS Position ^c
	11	12	13	14	15	16	17	18	19	20		
	G	G	G	G	G	G	G	G	G	G	1	3591
	C	C	C	C	C	C	C	C	C	C	2	3697
35	C	C	C	C	C	C	C	C	C	C	3	3804
	G	G	G	G	G	G	G	G	G	G	4	3818
	C	C	C	C	C	C	C	C	C	C	5	4123
	G	G	G	G	G	G	G	G	G	G	6	4240
	G	G	G	G	G	G	G	G	G	G	7	4472
40	G	G	G	G	G	G	G	G	G	G	8	4499
	A	G	G	G	G	G	G	G	G	G	9	4531
	G	G	G	G	G	G	G	G	G	G	10	4574
	C	C	C	C	C	C	C	C	C	C	11	4736
	T	C	C	C	C	C	C	C	C	C	12	4813
45	C	C	C	C	C	C	C	C	T	T	13	5068
	G	G	G	G	G	G	G	G	G	G	14	5103
	G	A	G	G	G	G	G	G	G	G	15	5150
	G	G	G	G	G	G	G	G	A	G	16	5179
	G	G	G	G	G	G	G	G	G	G	17	5301
50	G	G	A	G	G	G	G	G	G	G	18	5333
	G	G	G	C	G	G	G	G	G	G	19	5448
	G	G	G	G	A	G	G	G	G	G	20	5560
	G	G	G	G	G	A	G	G	G	G	21	5580
	T	C	C	C	C	C	C	T	C	C	22	5587
55	G	G	G	G	G	G	G	G	G	G	23	5606

	Haplotype Number ^a				PS ^b	PS
	21	22	23	24	Number	Position ^c
60	G	G	G	G	1	3591
	C	C	C	T	2	3697
	C	C	T	C	3	3804
	G	G	G	A	4	3818
	C	T	C	C	5	4123
	G	G	G	G	6	4240
	G	G	G	G	7	4472
65	G	G	G	G	8	4499
	G	G	G	G	9	4531
	T	G	G	G	10	4574
	C	C	C	C	11	4736
	C	C	C	C	12	4813
	C	C	C	C	13	5068
	G	G	G	T	14	5103
70	G	G	G	G	15	5150
	G	G	G	G	16	5179
	G	G	G	G	17	5301
	G	G	G	G	18	5333
	G	G	G	G	19	5448
	G	G	G	G	20	5560
	G	G	G	G	21	5580
75	C	C	T	C	22	5587
	G	G	G	G	23	5606
80						

^aAlleles for haplotypes are presented 5' to 3' in each column

^bPS = polymorphic site;

^cPosition of PS in SEQ ID NO:1.

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POLYMORPHISMS IN THE EDG6 GENE

ACGCCTGTGT	TCTCATGGGA	CGCCTGTGTT	CTCATGGGAC	GCCTGTGCC	
CTCATGGGAC	GCCTGTGTC	TCATGGGACG	CCTGTGCC	CATGGGACGC	100
CTGTGTTCTC	ATGGGACGCC	TGTGCCCTC	ATGGGACGCC	TGTGTTCTCA	
TGGGACGCC	GTGCCCTCA	TGGGACACCT	GTGCCCTCA	GGGACGCC	200
TGCCCTCATG	GGACGCC	GTTCTCATGG	GACGCC	CCCCTCATGG	
GACGCC	CCCCTCATGG	ACGCC	TCTCATGG	CGCCTGTGCC	300
CCTCATGGGA	CACCTGTGTT	CTCATGGGAC	GCCTGTGCC	TCATGGGACG	
CCTGTGTTCT	CATGGGACGC	CTGTGCCCT	CATGGGACAC	CTGTGTTCTC	400
ATGGGCCCTC	ATGGGACACC	TGTGTTCTCA	TGGGACGCC	GTGCCCTCA	
GGGACGCC	TGTGTTCTCA	GGACGCC	GCCCCTCA	GGACACCTGT	500
GCCCTCATGG	GACGCC	CCCCTCATGG	ACGCC	TCTCATGGGA	
CGCCTGTGCC	CCTCATGGGA	CGCCTGTGTT	CTCATGGGAC	GCCTGTGCC	600
CTCATGGGAC	ACCTGTGTC	TCATGGGACG	CCTGTGCC	CATGGGACGC	
CTGCGTTCTC	ATGGGACGCC	TGCATTCTCA	TGGGACGCC	GTGCCCTCA	700
GGGACACCTG	TGTGTTCTCA	GGACACCTGT	GTTCTCATGG	GATGCC	
CCCTCATGGG	ATGCC	CCCTCATGG	ACGCC	TCTCATGGGA	800
TGCC	TAC	CCCTCATGG	GGAC	ATCTGTGCTC	
TCATGGGATG	CCTGTGCC	TCATGGGATG	CCTGTGCC	CATGGGACGC	900
CTGCATTCTC	ATGGGACACC	TGTGCCCTCA	TGGGATGC	GTACCC	
GGGATGCC	TAC	GGACGCC	GCCC	GGATGCC	1000
GTTCTCATGG	GATGCC	CCCCTCATGG	GACGCC	TTCTCATGGG	
ACACCTGTG	CCTCATGGGA	TGCCTGTACC	CTTCATGGGA	CGCCTGTG	1100
TGGTTGCCAT	GATTACTACC	TGAGACTGTC	ACTACGACAG	TTACTATTGT	
TACTACTTGA	GACCATCATT	ACAAGACTGA	ACGAAGGGAC	GAATGTAGAA	1200
ATGAAA	AAGACAGAAG	AAACTGTTT	AAAGGAAGGG	ACCAGGGGAA	
GAAAAAAGAGA	GCTCC	TCTAGTGAGC	AAAGGCAGCC	CCCCAAGCTT	1300
CTACAGCC	TCG	TGGTAGAAA	GCAGGGGAG	GAAACGATTG	
GCCAGCTG	TGATTGTTCA	CACGTTCACG	TTATTGCTAA	CAGGTTTCAG	1400
ATTTGCCTAC	TTGCAAGAAA	CACTTG	TGGGGCGTGA	CTGCC	
CATTCC	GGGCGG	CAGCTTGT	CAGTTGCCA	ACAGC	1500
TTCATGAGAA	CAGTTG	TTTACTCACG	TAGCCTCCAG	TGGTATA	
AGTTGATCAC	AACCCTCATT	CTTCGGC	TCAACACCTG	AGCC	1600
GGACATCTG	GCCC	GGACACCTGT	GTCCTCGCAG	TACACT	
ACCC	GGG	GGGAGGAG	GCAGG	GAAACGATTG	1700
TGGTAGAATT	AGTGTAGCTG	CCCCCACC	CCCCCACC		
GAGGCCAAGG	ACACCATTG	CTCAGGAAGG	CTGAAGACCA	CAGGCTCCTG	
GGGGGACAGA	GGGCAAGG	GGCC	GACC	GGTGGTAAGT	1800
GGGC	TGGGGGTGAT	TGCAGGC	AGGAGG	CAGCAGGGAC	
TTATCCTGGG	TCTACTCAC	ACTTCTGGG	CCTGCATTAT	TTCCC	1900
ACCCCACACC	CCAAGGC	CTGGATGGG	ACGAGTGGG	GGTC	
ACTGGGGGAG	CTGGAGAGCA	GAGACCTCAC	ACTCCATCCG	TGACAGA	2000
TGTCCAAGCC	CCTACATGCC	CCAGAC	GGGCAAGG	GAGC	
CCTCAGACCC	CAGGGCAAGG	CTGAGCATCC	CCACTCAGAC	CTCAGGG	2100
GGCCTCGC	CCTCC	ACCCCAGGAC	AGGGC	CTCTCCC	
AGACCCCAAG	GCAGGGC	GCCTCCCC	TCAGA	GACAGGGCA	2200
TGCC	CTCAGACCC	AAGGCAAGG	CAAC	CCCTAGACCC	
CAGGCAGG	CAAGC	CGCTCAAAC	TCAGGG	GCATAC	2300
CTCCTCAGAC	CCAGGGC	GTGTG	CCTCTCAGAC	CACAGC	
GGCCTCGC	CCCTC	ACCCCAGGAC	AAGGCTGAGC	CTCTCCG	2400
AGACCCCAAGG	GTAAGGT	GCCTCCC	TCAGAAC	GGGCAAGGCC	
AACCC	CCTCAGACCC	CAGGCAGG	CAAGCTT	CCTCAGACGC	2500
CAGGGCAGAG	CCTGC	TCCTCAGACG	CCAGAGC	GTGTG	
CACTTAGACC	CACAGCC	TCACCTCAGG	CTGAGTC	ACC	2600

FIGURE 1A

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GTCAGCAGGG	CCATGGGGAT	GGGGTGAGCA	AGTCCCTACT	TTTCCTATGC	
ACCTCAATCC	CAGTGGGGGG	CTGCCCCAGG	GGGCCAGCAG	CTCTGCTCCC	2700
AGCAGGGTGA	GCTCAGGGGC	AGGAGAGGCA	ACCATGAATC	CCAAAATGGG	
CGCCTGGCCC	CGAGACCCTA	CCAGCTTGTG	CCTGGGGTC	TCTCTCCCTG	2800
GACCTGTGTG	TCCTCCCATG	GGCAGAAAGT	TGGCCTCAGG	CCGCCTTGT	
GGGCCTCAGT	TTTCCCTTCT	GTAACGCATC	AGGCTCTCTG	GGCCTCAGTT	2900
TCCCCCTTCTG	CAACGCATCG	GGCTCTCTGG	GCCTCAGTTT	CCCCCTCTGC	
AACGCATCGG	GCTCTCTGGG	CCTCAGTTTC	CCCTTCTGCA	ACGCATCGGG	3000
CTCTCTGGGC	CTCAGTTTCC	CCTTCTGCAA	CGCCTCGGGC	TCTCTGGGCC	
TCAGTTTCCC	CTTCTGCAAC	GCATCGGGCT	CTCTGGGCCT	CAGTTTCCCC	3100
TTCTGCAACG	CCTCGGGCTC	TCTGGGCCTC	AGTTTCCCCT	TCTGCAACGC	
CTCAGGGCTCT	CTGGGCTCTA	GTTCCTCCCTT	CTGCAACGCA	TCAGGGCTCTC	3200
TGGGCCTCAG	TTTCCCTCTC	TGCAACGCAT	CGGGCTCTCT	GGGCCTCAGT	
TTCCCCTTCT	GCAACGCATC	GGGCTCTCTG	GGCCTCAGTT	TCCCCCTCTG	3300
CAACGCCTCG	GGCTCTCTGG	GCCTCAGTTT	CCCCCTCTGC	ACAGCCTCGG	
GCTCTCTGGG	CCTCAGTTTC	CCCTTCTGCA	ACGCATCGGG	CTCTCTGGGC	3400
CTCAGTTTCC	CCTTCTGCAA	CGCATCGGGC	TCTCTGGGCC	TCAGTTTCCC	
CTTCTGCAAC	GCACCGGGCT	CTCTGGGCCT	CAGTTTCCCTC	TTCTGTAACG	3500
CATCAGGGCCC	TGTTTCGGGG	ATCAAGTCGG	ATGAGTCAGT	GCTCAAGGGC	
ATGCAGGCAC	TTGACATTAA	TTAGGCACCT	GCTGTGTGCT	GAGCGCCGGT	3600
			A		
GGAGCATGTG	GGGAGACCTC	AGTGGAGCCG	ACGGGTGCTT	CGGGGGTGAT	
CAGGGCTGGT	GGGGAGGGAGT	CGTCCGGCTG	GAAAGGGGTG	GCCCCATCCCG	3700
			T		
AGTGGGGACT	CATTTCCCT	CCGTGACTGA	CGGCTCCGGG	GCTCCCTGCG	
GGTCTCCGGC	TTTCGAGGAC	AGGAAGAAGG	CAGCCAGGGC	AGGGGTGGGG	3800
GTCCTCACAG	CCAGGGCAGC	CCCAGCGCGT	TGGCTCCAGG	AGCCCAGGGTG	
		G			
GGGGCCGACC	GTTGGGGTGC	CCCTCCCTGT	CCTCGGCCTT	ACCTCCACCC	3900
TGGAGGGCAC	CTTGAACATA	ACAGGAAATT	TCAAATAACA	GGAAACCAAAG	
ACCAGCAAGG	CGGGTGGCTC	CACCCCTGCGT	CGGGCCTGAG	TCAGCCCCCG	4000
GGGGAGGGCCA	TGAACGCCAC	GGGGACCCCCG	GTGGCCCCCG	AGTCCTGCCA	
	[exon 1: 4010..				
ACAGCTGGCG	GCCGGCGGGC	ACAGCCGGCT	CATTGTTCTG	CACTACAACC	4100
ACTCGGGCCG	GCTGGCCGGG	CGCGGGGGGC	CGGAGGATGG	CGGCCTGGGG	
		T			
GCCCTGCGGG	GGCTGTCGGT	GGCCGCCAGC	TGCCTGGTGG	TGCTGGAGAA	4200
CTTGCTGGTG	CTGGCGGCCA	TCACCAGCCA	CATGCGGTG	CGACGCTGGG	
		A			
TCTACTATTG	CCTGGTGAAC	ATCACGCTGA	GTGACCTGCT	CACGGCGCG	4300
GCCTACCTGG	CCAACGTGCT	GCTGTCGGGG	GCCCCACCT	TCCGTCTGGC	
GCCCCCCCAG	TGGTTCTAC	GGGAGGGCCT	GCTCTTCACC	GCCCTGGCCG	4400
CCTCCACCTT	CAGCCTGCTC	TTCACTGAG	GGGAGCGCTT	TGCCACCATG	
GTGCAGGCCGG	TGGCCGAGAG	CGGGGCCACC	AAGACCAGCC	GCGTCTACGG	4500
		A	A		
CTTCATCGGC	CTCTGCTGGC	TGCTGGCCGC	GCTGCTGGGG	ATGCTGCCTT	
		A			
TGCTGGGCTG	GAACTGCCTG	TGCGCCTTTG	ACCGCTGCTC	CAGCCTTCTG	4600
		T			
CCCCCTACT	CCAAGCGCTA	CATCCTCTTC	TGCCTGGTGA	TCTTCGCCGG	
CGTCCTGGCC	ACCATCATGG	GCCTCTATGG	GGCCATCTTC	CGCCTGGTGC	4700
AGGCCAGCGG	GCAGAAGGCC	CCACGCCAG	CGGCCCGCCG	CAAGGCCCGC	
		T			
CGCCTGCTGA	AGACGGTGCT	GATGATCCTG	CTGGCCTTCC	TGGTGTGCTG	4800

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GGGCCCACTC	TTCGGGCTGC	TGCTGGCCGA	CGTCTTGCG	TCCAACCTCT	
T					
GGGCCAGGA	GTACCTGCAG	GGCATGGACT	GGATCCTGGC	CCTGGCCGTC	4900
CTCAACTCGG	CGGTCAACCC	CATCATCTAC	TCCTTCCGCA	GCAGGGAGGT	
GTGCAGAGCC	GTGCTCAGCT	TCCTCTGCTG	CGGGTGTCTC	CGGCTGGGCA	5000
TGCGAGGGCC	CGGGGACTGC	CTGGCCCCGG	CCGTCGAGGC	TCACTCCGGA	
GCTTCCACCA	CCGACAGCTC	TCTGAGGCCA	AGGGACAGCT	TTCGCGGCTC	5100
T					
CCGCTCGCTC	AGCTTTCGGA	TGCGGGAGCC	CCTGTCCAGC	ATCTCCAGCG	
T				A	
TGCGGAGCAT	CTGAAGTTGC	AGTCTTGCCT	GTGGATGGTG	CAGCCACCGG	5200
A					
..5164]					
GTGCGTGCCA	GGCAGGGCCT	CCTGGGGTAC	AGGAAGCTGT	GTGCACGCAG	
CCTCGCCTGT	ATGGGGAGCA	GGGAACGGGA	CAGGCCCTCA	TGGTCTCCC	5300
GGTGGCCTCT	CGGGGCTTCT	GACGCCAAAT	GGGCTTCCA	TGGTCACCCT	
A		A			
GGACAAGGGAG	GTAACCACCC	CACCTCCCCG	TAGGAGCAGA	GAGCACCCCTG	5400
GTGTGGGGGC	GAGTGGTTCC	CCACAACCCC	GCTTCTGTGT	GATTCTGGGG	
C					
AAGTCCCGGC	CCCTCTCTGG	GCCTCAGTAG	GGCTCCCAGG	CTGCAAGGGGG	5500
TGGACTGTGG	GATGCATGCC	CTGGCAACAT	TGAAGTTCGA	TCATGGTACG	
TGATGTTGCG	GCCTCTTATT	CCCTGGTGCG	TGCATGCGTG	GGGGCCGTGG	5600
A		A	T		
CTCAGGGGGG	CTGTGGATCT	AGGGGCAGCC	GGGTGTGTCT	TTGCTAGAGA	
C					
GGGCCACGGG	CCAGTGCCT	GTGAGGGTGG	AGTGTGTGTG	TGTGTGTGTG	5700
TGTGTGTGTG	TGTGTGTGGA	CAACCTCTGG	GCCTTGCGGG	AAGTGGGGGT	
GACAATGACA	TTAAATGCGC	CCTCTTCTTG	TTCACCTCCC	CTTTAGAAAT	5800
GGCAGGGCCC	ATGCCCATC	TCTGGCCTCT	GCATCTTTG	GGGACCCACT	
CTCTGGGGCT	GGCAGAGGCA	CCACCTTGGC	TTCCTGGGCT	GGGGGAATCT	5900
TCCCCTCACAT	CCCCTTCAGC	ATGAACGGCC	TCGGCTTCC	CGGTGGTAA	
AACAGTTAA	TCACTGAAGC	CGAACGACAG	GGTTGATTGT	ACACGCTCCC	6000
CGCCAGGCCAC	AGGGGCTGAC	AACTGCCTGC	CCCCTGAAAC	TCCAGTGGAG	
ACGTTTCAGC	TCCACACCAT	TCAGTATGGG	AGACGCCAGC	CCCACGGGC	6100
TACGGTGCAA	GCAGATAACT	GAATTCGAA	GTGTAGGTTG	TGTTTAATTT	
GAATCTGTTT	ATATTTGGT	AGCCCCATGG	GGCGGGTGGC	CACAGTTCA	6200
GTGCAGATGT	AAATCCGAA	GCCTCCAGCA	CCTGCAGCTC	ATAGACAGCT	
CTCGCCCCACC	TTCTCCCAGG	ACCAAGCCAG	TCCCCTCCAG	TCCAGTGTCT	6300
GAGCAGAGTC	AGAATCCACA	CCACCCGCCG	CCTGGGCTCA	GAAAGTCTG	
CTTTAAGTCA	TTATTTCTCC	ACTGTACGAT	GGGAAATGCG	GTGTGTGGGG	6400
GCCATTCTACC	CACCAACACA	GCAGCTGTGA	GGCACACACG	GCTATTGAAA	
ATTCATGGAA	ATTGCTGGGT	GTGGTGGCTC	ATGCCTGTAA	TCCCAGCACT	6500
TTGGGAGGCC	GAGGCAGGAG	GATTGCTTGA	GTGCAGGAGT	TCCAGACCAAG	
CCTGGGCAAC	ATAGCGAAAC	CACATCTCTA	CAAAAAAAATC	CTCCAAAATT	6600
AAAAAAATTA	GCCCGAGCAT	GTTGTTGCAT	GCCTGTGGTT	CCAGCTACTC	
GAGAGGCTGA	GGTGGGAGGA	TCACTTGAGC	CCGGGAGGTC	GAGGCTGCAG	6700
GGAGCGGTGA	TCGTGCCAGC	CTGGGTGACA	GAGTGAGACC	CTGTCTCTAA	
AAAAAAATTA	AAAAAAATTA	AAAAAGAAAA	TTCATTGAAA	TTAAATGAAA	6800
GTTAGGCCAG	GTGCATTGGT	TCATGCCTGT	AATCGCAGCA	CTTTGGGGGG	
CCGAGGTGGG	TGGATCATGA	GGTCAGGAGT	TCAAGACCAG	CCTGGCTAAG	6900
ATGGTGAACAC	CCTGTCTCTA	CTAAAAATAC	AAAAAATAAG	CTGGGTGCAG	
TGGCAGGGCGC	CTGTAATCCC	AGCTACTCGG	GAGGCTGAGG	CAGGAGAATC	7000
GCTTGAACTC	GGAGGGTGGG	GGTGCAGTG	AGCTGGGATT	GCTCCACTGC	

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ACTCCAGTCT	GGCGGATAGA	GTGAGACTCC	GTCTCAAAAA	ACAAAACAAA	7100
AACAACAACA	AAAAAGAAAT	TAAATGAAAG	TTAAAAAATT	AGTCTCTCAG	
TGTATTGATT	TGCTAGGGCA	GCCATGACAG	TCTCACAGAT	GGAGAGGCTT	7200
GAAAACAGAC	ATTTATGCTG	CCATAATTCT	GGAAGCCAGA	AGTCTGAGAT	
GAAGGTGTGA	GTAGGGCTGG	GTCCTCCTGA	GGCTGACGGG	GATCTGCCA	7300
GGCCTCTCCC	AGTCCTGCT	GGTGGCCGGC	GTCCTTGATG	TTCCCTGGCT	
TGGAGAACG	TCAGCCGCAT	CTCTGCCCTCC	ATCTCCACAA	GGCGCCCTCC	7400
CTGGGTCTGT	ATCCAAACTC	CCCCAACTCC	TTCAACCTTT	TTTATTTAA	
GAGATGAGGG	GCCGGGCGCC	GTGGCTCACG	CCTGTAATCC	CAGCACTTTG	7500
GGAGGCCGAG	GTGGGCGGAT	CACGAGGTCA	GGAGATAGAG	ACCATCCTGG	
CTAACACGGT	GAAGCCCCAT	CTTTACTAAA	AATACAAAGA	ATTAGCCAGG	7600
CGTGGTGGCG	GGTGCCTGTA	GTCCCAGCTA	CTCTGGAGGC	TGAGTCAGGA	
GAATGGCGTG	AACCCGGGAG	GCAGAACTTG	CAGTGAGCGG	AGATCAGGCC	7700
ACTGCACTCC	ACCCCTGGGTG	ACAGAGAGAG	ACTCCGTCTC	AAAAAAAAAAA	
AAAAAAAAAAA	AAAAAAAAGA	GATGAGGGTC	TCATTAAGTT	GCCCCAGGCTG	7800
GTCTCAAAC	CCTGGGCTCA	AGTGATCCTC	CCACCTCAGC	CTTCTGAATA	
GCTGAGACTA	CAGGAGTGCA	CCACCAAGCC	TGGCTCATTT	TCCTATTTT	7900
ACATTATTAA	TTATTATTGT	TATTATTATT	TTTTTGAGAA	GGACTCTCGC	
TCTGTCGTCC	AGGCTGGAGC	GCAGTGGCGC	GATCTCAGCT	CACCACAACC	8000
TCCACCTCCT	GGGTTCAAGC	GATTCTCCTG	CCTCAGCCTC	CCGACTGGGA	
CTACAGGCAC	GCACCATCAT	ATCTGGCTAG	TTTTTGTATT	TTTAGTAGAG	8100
ACGGGATT	GCCATGTTGG	CCAGGCTGGT	CTCGAACTCC	TGACTCAGGT	
GATCCATCTG	CCTCCCAAAG	TGCTGGGATT	ACAGGCGTGA	GCCACCGTGC	8200
CCGGCCGATT	TCCCCATTTT	TATAAGGCCA	CCAGTCACAT	GGGATTAAGG	
GCCCACCCCTG	CTCCAGTATG	ACCTCATCTT	AACTAATGCC	ATCTGCCACA	8300
ACCATTTCC	AAATAAACATC	CTATGCTGAG	GTCCTGGGGG	TTAGGATGTG	
ATCATGTAGA	TTTGATGGG	ACAGTTCTTT	ATTTTCTTTT	ATATTTAGA	8400
GACAGGGTCT	TGCTCTGTCA	CCCAGGCTGG	AGTGCAGTGG	TGTGATCATA	
GCTCACTGCA	GCCTCGATCT	CTTGGGCTCA	AGTGATCCTT	CTGCCTTGGG	8500
TTTCCGAGTA	GCTGCAATT	CAGGTGCACA	CCACCACACC	TGGCTAATTT	
TTATATTTT	TGTAGAGACG	GGGCTGGGTG	CTGTGGCTCA	CCCCTGTAAT	8600
TCCAGCACTT	TGGGAGGCCG	AGGCAGGGTGG	ATCACCTTAG	GTCAGGAGTT	
TGAGACCAGC	CTGGCCAACA	TGGAGAAACC	CCGTCTCTAC	AAAAAATACA	8700
AAATTAGCCA	GGCGTAGTGG	TGCATGCCTG	TAATCCCAGC	TACTTGGGAG	
GCTGAGGCAG	GAGAATTGCT	TGAACCCGGG	AGGCCGAGGT	TGCAGTGGAC	8800
TGAGTTCGTG	CCACTGCACT	CCAATCTGGG	TGACAGAGCA	AGACTCCGTC	
TCAAAAAAAA	AAAAAAAAGA	AAAAGAGAGA	GAGAGAGAGA	GACAGGGTCT	8900
TGCTATGTT	CCCAGGCTGG	TCTCAATCTC	CTGGGCTCAA	GTGATCCTCC	
TGCCTCAGCC	TCCCCAAAGTG	TTGGGACTAC	AGGCATGAGC	TACTGTGGCT	9000
GGCCAATTGC	AGACAGATGG	AACTGTGGCC	AGCCACTTCA	CCAGTGAAGT	
CCCAAACACC	CCTCTGCAGG	CATGATAAGC	TAGGGTAGTT	CTTATCTTT	9100
CCTGATTCC	GTTCTCTGTG	TGTTTAGTGG	GGACCACAGC	AGGCAGGGTG	
GGGTGTTGGT	GGA				9163

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POLYMORPHISMS IN THE CODING SEQUENCE OF EDG6

ATGAACGCCA CGGGGACCCC GGTGGCCCCC GAGTCCTGCC AACAGCTGGC GGCCGGCGGG CACAGCCGGC TCATTGTTCT GCACTACAAC CACTCGGGCC GGCTGGCCGG GCGCGGGGGG CCGGAGGATG GCAGGCCTGGG GGCCCTGCGG T	100
GGGCTGTCGG TGGCCGCCAG CTGCCCTGGTG GTGCTGGAGA ACTTGCTGGT GCTGGCGGCC ATCACCAGCC ACATGCGGTC GCGACGCTGG GTCTACTATT A	200
GCCTGGTGAA CATCACGCTG AGTGACCTGC TCACGGGCGC GGCCTACCTG GCCAACGTGC TGCTGTCGGG GGCCCACCGC TTCCGCTCTGG CGCCCGCCCA GTGGTTCTA CGGGAGGGCC TGCTCTTCAC CGCCCTGGCC GCCTCCACCT TCAGCCTGCT CTTCACTGCA GGGGAGCGCT TTGCCACCAT GGTGCGGCCG GTGGCCGAGA GCGGGGCCAC CAAGACCAGC CGCGTCTACG GCTTCATCGG A A	300
CCTCTGCTGG CTGCTGGCCG CGCTGCTGGG GATGCTGCCT TTGCTGGGCT A	400
GGAACTGCCT GTGCGCCTTT GACCGCTGCT CCAGCCTTCT GCCCCCTCTAC T	500
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AAGACGGTGC TGATGATCCT GCTGGCCTTC CTGGTGTGCT GGGGCCCACT CTTCGGGCTG CTGCTGGCCG ACGTCTTGG CTCCAACCTC TGGGCCAGG T	700
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 nnnnnnnnnn nnnnnnnnnn nnnnnnnnnn nnnnnnnnnn nnnnnnnnnn 2760